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# The HIFI spectral survey of AFGL 2591 (CHESS)

## II. Summary of the survey<sup>★,★★</sup>

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### ABSTRACT

**Aims.** This paper presents the richness of submillimeter spectral features in the high-mass star forming region AFGL 2591.

**Methods.** As part of the Chemical *Herschel* Survey of Star Forming Regions (CHESS) key programme, AFGL 2591 was observed by the *Herschel* (HIFI) instrument. The spectral survey covered a frequency range from 480 to 1240 GHz as well as single lines from 1267 to 1901 GHz (i.e. CO, HCl, NH<sub>3</sub>, OH, and [CII]). Rotational and population diagram methods were used to calculate column densities, excitation temperatures, and the emission extents of the observed molecules associated with AFGL 2591. The analysis was supplemented with several lines from ground-based JCMT spectra.

**Results.** From the HIFI spectral survey analysis a total of 32 species were identified (including isotopologues). Although the lines are mostly quite weak ( $\int T_{\text{mb}} dV \sim \text{few K km s}^{-1}$ ), 268 emission and 16 absorption lines were found (excluding blends). Molecular column densities range from  $6 \times 10^{11}$  to  $1 \times 10^{19} \text{ cm}^{-2}$  and excitation temperatures from 19 to 175 K. Cold (e.g. HCN, H<sub>2</sub>S, and NH<sub>3</sub> with temperatures below 70 K) and warm species (e.g. CH<sub>3</sub>OH, SO<sub>2</sub>) in the protostellar envelope can be distinguished.

**Key words.** ISM: individual objects: AFGL 2591 – ISM: molecules – stars: formation – stars: massive

## 1. Introduction

Massive stars play a major role in the evolution of galaxies. From their birth in dense molecular clouds to their death as a supernova explosion, massive stars interact heavily with their surroundings by emitting strong stellar winds and by creating heavy elements (Zinnecker & Yorke 2007). They influence the formation of nearby low-mass stars and planets (Bally et al. 2005) and the physical, chemical, and morphological structure of galaxies (e.g. Kennicutt & Evans 2012). Although massive stars are an important component of galaxies, their formation processes are still unclear. It is difficult to observe high-mass star forming regions because of high dust extinction and their large distances and rapid evolution (Tan et al. 2014).

High-mass star forming regions are quite rare, so each observational effort is very helpful in solving they represent. One of the goals of the *Herschel* Space Observatory (Pilbratt et al. 2010) was to improve our understanding of the high-mass star formation processes. Among the several key projects devoted to those studies, we focus here on the *Herschel* key program Chemical *Herschel* Survey of Star Forming Regions (CHESS, Ceccarelli et al. 2010). The aim of this project is to study the chemical composition of dense regions of the interstellar medium and to understand the chemical evolution of star

forming regions and the differences between regions with different masses/luminosities. The target sources of CHESS are the pre-stellar cores I16293E and L1544, the outflow shock spot L1157-B1, the low-mass protostar IRAS16293-2422, the intermediate-mass protostar OMC2-FIR 4, the intermediate-luminosity hot cores NGC 6334I and AFGL 2591, and the high-luminosity hot core W51e1/e2. Almost the entire spectral range of the HIFI instrument, that is, 480 to 1910 GHz, has been used to observe these objects. In this paper we focus on the source AFGL 2591.

Spectral surveys simultaneously cover a wide variety of molecular and atomic lines. In this way they offer the possibility to probe cold and warm gas and the fundamental processes that occur in star forming regions. Especially the wide frequency range of *Herschel*'s allowed us to cover molecular lines from very different energy levels, from light to heavier molecules, and thus to study the several species thoroughly.

AFGL 2591 is one of the CHESS sources. It is a relatively isolated high-mass protostellar object with a bipolar molecular outflow (Van der Tak et al. 1999). A massive sub-Keplerian disk has been proposed to exist around source AFGL 2591–VLA 3 (Wang et al. 2012). AFGL 2591 is located in the Cygnus X region, ( $l, b$ ) = 78°9, 0°71. Based on VLBI parallax measurements of 22 GHz water maser, Rygl et al. (2012) have recently estimated the distance<sup>1</sup> towards AFGL 2591 to be  $3.33 \pm 0.11 \text{ kpc}$ ,

<sup>★</sup> *Herschel* is an ESA space observatory with science instruments provided by European-led Principal Investigator consortia and with important participation from NASA.

<sup>★★</sup> Appendix A is available in electronic form at <http://www.aanda.org>

<sup>1</sup> The previous distance estimates were uncertain, with values between 1 and 2 kpc (e.g. Van der Tak et al. 1999, 2000b), thus luminosity at 1 kpc  $L = 2 \times 10^4 L_{\odot}$ .

hence, the corresponding luminosity is  $L = 2 \times 10^5 L_{\odot}$  (Sanna et al. 2012). For a detailed source description see Van der Wiel et al. (2013, hereafter Paper I) and references therein.

The richness of the detected lines in AFGL 2591 from the HIFI/CHESS spectral survey gives us the opportunity to gain detailed insight into its chemical and physical structure. Results from the spectral survey will be presented in a series of papers. The first one focused on highly excited linear rotor molecules (Van der Wiel et al. 2013). In the present work the entire HIFI spectral survey of AFGL 2591 is presented.

Van der Wiel et al. (2013) studied linear rotor molecules (CO, HCO<sup>+</sup>, CS, HCN, HNC) in the high-mass protostellar envelope. This work was based on the *Herschel*/HIFI data together with observations from the ground-based telescopes, JCMT and IRAM 30 m. The line profiles of the observed emissions consist of two components, a narrow one that corresponds to the envelope, and a broad component from the outflow. The same nomenclature is used in the present paper.

This paper starts with the description of the observations and the data reduction of *Herschel* and JCMT spectra (Sect. 2). In Sect. 3 the general summary of the HIFI/CHESS spectral survey of AFGL 2591 is given. Here, all of the observed species from that survey are presented together with emission and absorption lines analysis. Discussions and conclusions are given in Sects. 5 and 6. Appendix A gives a table with all detected transitions and plots of their line profiles.

## 2. Observations and data reduction

### 2.1. 480–1850 GHz *Herschel*/HIFI data

Observations of AFGL 2591 ( $\alpha_{2000} = 20^{\text{h}}29^{\text{m}}24^{\text{s}}.9$ ,  $\delta_{2000} = +40^{\circ}11'21''$ ) were obtained with the Heterodyne Instrument for the Far-Infrared (HIFI, de Graauw et al. 2010) on board the ESA *Herschel* Space Observatory as a part of the HIFI/CHESS guaranteed time key programme<sup>2</sup>.

A full spectral survey of AFGL 2591 of HIFI bands 1a–5a (480–1240 GHz, 18.4 h of observing time) was obtained. Nine additional selected frequencies were observed in 3.5 h of observing time. The corresponding bands are 5b (lines: HCl, CO), 6a (CO), 6b (CO), 7a (NH<sub>3</sub>, CO) and 7b (CO, OH, [CII]).

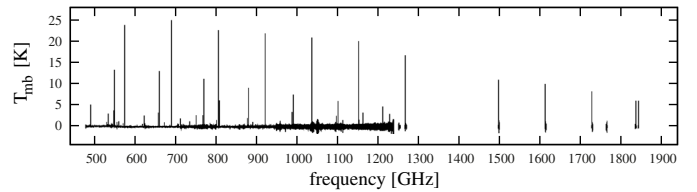
Although this is the second in a series of papers based on HIFI/CHESS data of AFGL 2591 and a detailed description of its data reduction process has been provided in Paper I (Van der Wiel et al. 2013), basic information is recalled here as well.

The spectral scan observations were carried out using the dual beam-switch (DBS) mode, with the Wide Band Spectrometer (WBS) with a resolution of 1.1 MHz, corresponding to 0.66 km s<sup>-1</sup> at 500 GHz and 0.18 km s<sup>-1</sup> at 1850 GHz. The single-frequency settings were obtained in the dual beam-switch mode as well, with the fast chop and stability optimization options selected. Table 1 gives information about the covered frequency range, beam size, noise level, and integration time.

AFGL 2591 data were completely reduced with the *Herschel* interactive processing environment<sup>3</sup> (HIPE; Ott 2010), version 8.1, using scripts written by the CHESS data reduction team (Kama et al. 2013). After pipelining, the quality of

**Table 1.** Overview of all HIFI bands for the observations of AFGL 2591 (bands 1a–5a are spectral scans, 5b–7b single-frequency settings).

Band	Freq. range [GHz]	Beam size [']	rms [K]	Obs. time [s]
1a	483–558	41	0.030	4591
1b	555–636	36	0.029	4643
2a	631–722	31	0.026	9833
2b	717–800	28	0.067	6407
3a	800–859	26	0.039	4893
3b	858–960	23	0.067	8578
4a	950–1060	21	0.157	9137
4b	1051–1120	20	0.144	6300
5a	1110–1240	18	0.147	11931
5b	1266–1270	17	0.149	1380
5b	1251–1255	17	0.149	2255
6a	1496–1499	14	0.117	1440
6b	1611–1614	13	0.106	1392
7a	1726–1729	12	0.092	1575
7a	1762–1764	12	0.095	1423
7b	1840–1843	12	0.092	1711
7b	1900–1903	12	0.120	1452



**Fig. 1.** Complete baseline-subtracted spectrum. The strongest lines belong to CO and its isotopologues, while at 1901 GHz [CII] is seen.

each spectrum was checked and spectral regions with spurious features (spurs) were flagged. Next, the correction for standing waves was made and a baseline was subtracted (polynomial of  $\sim 3$ ). The final single-sideband spectrum is presented in Fig. 1.

Strong lines are known to create *ghost* features in the side-band deconvolution process (Comito & Schilke 2002). To check the importance of this effect on our data, the above steps were repeated with strong lines (especially CO transitions) masked out in the same way as spurs. The term *strong lines* refers to features of  $T_A^* > 1$  K in band 1a to  $> 8$  K in band 5a, depending on the amount of lines and the noise level in a given band. Following the outlined data reduction procedure, two single-sideband spectra for bands 1a–5a were obtained. The first set of spectra was used to analyse strong lines (e.g. CO and its isotopologues, HCO<sup>+</sup>), the second for line measurements of weak features, that is, those that were not masked as strong lines (e.g. SO, CH<sub>3</sub>OH).

### 2.2. 330–373 GHz JCMT data

The excitation analysis of several molecules was complemented by ground-based observations from the *James Clerk Maxwell* Telescope (JCMT)<sup>4</sup>. These data are part of the JCMT Spectral Legacy Survey (SLS, Plume et al. 2007). The observations were taken with the 16-element Heterodyne Array Receiver Programme B (BHARP-B) and the Auto-Correlation Spectral

<sup>2</sup> Data are available from

[www-laog.obs.ujf-grenoble.fr/hebergues/hs3f](http://www-laog.obs.ujf-grenoble.fr/hebergues/hs3f)

<sup>3</sup> HIPE is a joint development by the *Herschel* Science Ground Segment Consortium, consisting of ESA, the NASA *Herschel* Science Center, and the HIFI, PACS and SPIRE consortia.

<sup>4</sup> The *James Clerk Maxwell* Telescope is operated by the Joint Astronomy Centre on behalf of the Science and Technology Facilities Council of the United Kingdom, the Netherlands Organisation for Scientific Research, and the National Research Council of Canada.

Imaging System (ACSIS) correlator (Dent et al. 2000; Smith et al. 2008; Buckle et al. 2009).

The JCMT survey of AFGL 2591 covers the frequency range of 330–373 GHz with a spectral resolution of 1 MHz ( $\sim 0.8 \text{ km s}^{-1}$ ). The beam size of the JCMT at these frequencies is  $14\text{--}15''$ , the image size is  $2'$ . Detailed information about the data reduction and analysis can be found in Van der Wiel et al. (2011).

### 3. HIFI spectral survey of AFGL 2591

#### 3.1. Detections and line profiles

From the *Herschel*/HIFI spectral survey, a total of 32 species (including isotopologues) were identified, resulting in 252 emission and 16 absorption lines (218 different transitions). Blended features were excluded from the analysis. *Herschel* surveys toward different sources revealed many spectral features that we currently cannot identify (e.g. Wang et al. 2011). However, no unidentified lines were found in our spectra.

For the line identification the JPL (Pickett et al. 1998) and CDMS (Müller et al. 2001, 2005) databases were used. Line analysis was made with the CASSIS software<sup>5</sup>. The presence of possible transitions resulting from an upper energy level  $E_{\text{up}}$  of less than 500 K was checked. Generally, detected lines have  $E_{\text{up}} < 400 \text{ K}$ , except for the high- $J$  CO transitions, which have  $E_{\text{up}}$  up to 752 K.

All the detected lines in the HIFI survey are presented in Table A.1, the entire spectrum is shown in Fig. 1, and corresponding line profiles can be found in Fig. A.1. For the sake of completeness all of the observed lines together with their profiles and measurements are presented in Table A.1, including the datasets of Paper I (Van der Wiel et al. 2013) and the complementary JCMT data.

Although the analysis of our survey revealed no new molecular species, some of our observed species have not been detected toward AFGL 2591 before. HIFI with its broad spectral range gave the opportunity to observe for the first time transitions of HF (Emprechtinger et al. 2012),  $\text{OH}^+$ , CH,  $\text{CH}^+$  (Bruderer et al. 2010b) or  $\text{C}^+$  and HCl (this work) in AFGL 2591.

Within the object AFGL 2591,  $\text{CH}_3\text{OH}$ ,  $\text{SO}_2$ , and SO show the highest number of detected transitions (54, 26, and 18 lines, respectively) among its identified species, followed by  $\text{H}_2\text{CO}$  and CO and its isotopologues. For the other molecules, a few lines at most were observed. The strongest transitions originate from CO and its isotopologues,  $\text{HCO}^+$ ,  $\text{H}_2\text{O}$ , and OH. In comparison, the remaining detected lines are relatively weak because of fluxes below  $1 \text{ K km s}^{-1}$ .

The lines were measured in the same way as described in Paper I. A Gaussian profile was fitted to each line, using the Levenberg-Marquardt fitter in the line analysis module of CASSIS. For most lines, a single Gaussian profile gave a good fit to the profile. However, for CO,  $^{13}\text{CO}$ ,  $\text{C}^{18}\text{O}$ , Cl, [CII],  $\text{HCO}^+$ , OH, and  $\text{H}_2\text{O}$ , double Gaussian profiles were needed to fit sufficiently narrow and broad line components. The measured parameters from Gaussian fits of the emission lines (central velocity and full width at half maximum) are plotted in Fig. 2 (together with the complementary JCMT data) as an average value for each molecule.

The narrow and single-line components are centred on  $-5.5 \pm 0.5 \text{ km s}^{-1}$  (as derived before by Van der Tak et al. 1999)

**Table 2.** Fit results for absorptions.

Molecule	$V_{\text{lsr}}^a$ [ $\text{km s}^{-1}$ ]	$\Delta V^a$ [ $\text{km s}^{-1}$ ]	$N$ [ $\text{cm}^{-2}$ ]
CCH	0.82	2.58	$3.3 \pm 1.0 \times 10^{17}$
$\text{CH}^b$	0.21	2.19	$3.1 \pm 0.9 \times 10^{13}$
$\text{CH}^+$	4.17	12.42	$1.1 \pm 0.4 \times 10^{14}$
$\text{H}_2\text{S}$	0.22	0.97	$3.5 \pm 0.9 \times 10^{12}$
$\text{NH}_3$	0.00	1.30	$1.8 \pm 0.8 \times 10^{12}$
$\text{H}_2\text{O}^b$	-0.50	2.43	$1.5 \pm 0.6 \times 10^{13}$
$\text{OH}^{+b}$	3.65	9.13	$3.0 \pm 1.0 \times 10^{13}$
HF	-0.05	2.31	$5.2 \pm 1.3 \times 10^{12}$
HF	-3.88	2.50	$5.5 \pm 1.4 \times 10^{12}$
$\text{H}_2\text{O}$	-11.98	13.75	$2.3 \pm 0.6 \times 10^{13}$
$\text{CH}^+$	-16.90	9.24	$6.8 \pm 1.3 \times 10^{13}$
HF	-12.58	8.81	$1.8 \pm 0.6 \times 10^{13}$

**Notes.** <sup>(a)</sup> Errors of  $V_{\text{lsr}}$  and  $\Delta V$  are listed in Table A.1. <sup>(b)</sup> The average of a few lines from the same velocity component: 2 lines of CH, 2 lines of  $\text{H}_2\text{O}$  and 3 lines of  $\text{OH}^+$ .

and originate from the protostellar envelope. Their line widths are of the order of  $3.7 \pm 0.9 \text{ km s}^{-1}$ , whereas the broader line components ( $10.9 \pm 4.2 \text{ km s}^{-1}$ ) are caused by the outflows and are centred on  $-6.3 \pm 0.7 \text{ km s}^{-1}$ . It was shown in Paper I that the outflow gas is not significantly different from that in the envelope, considering gas density, gas temperature, and the chemical balance of CO and  $\text{HCO}^+$ .

#### 3.2. Absorption line analysis

There are only a few absorption features observed toward AFGL 2591. A foreground cloud at  $V_{\text{lsr}} \sim 0 \text{ km s}^{-1}$  has been detected before, for example, Bruderer et al. (2010b), Emprechtinger et al. (2012), and Van der Wiel et al. (2013). In the CHESS/HIFI dataset we found 16 absorption lines; all measurements are listed together with emissions in Table A.1, and their lines profiles are presented in Fig. A.1. Most of them are red-shifted and associated with the foreground cloud at  $V_{\text{lsr}} \sim 0 \text{ km s}^{-1}$ . Three broad, blue-shifted absorptions belong to the outflow lobe.

We derived the molecular column densities using the following relations:

$$N_{\text{tot}} = N_1 \frac{Q(T_{\text{ex}})}{g_1} \exp\left(\frac{E_1}{kT_{\text{ex}}}\right) [\text{cm}^{-2}] \quad \text{and} \quad (1)$$

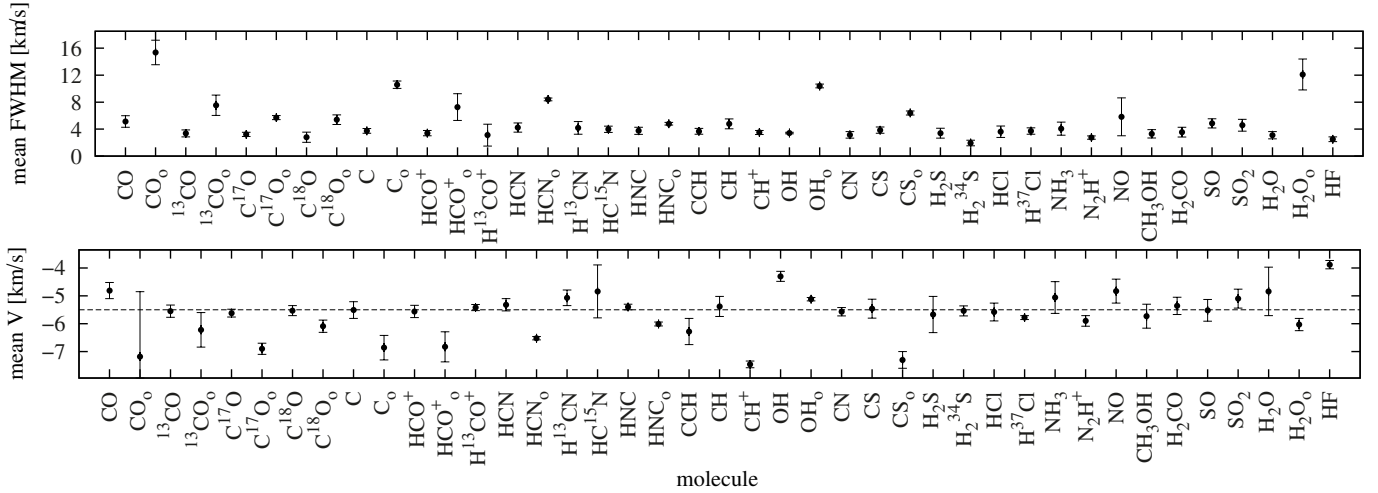
$$N_1 = \frac{8\pi\nu^3}{c^3} \frac{g_1}{g_u A_{ul}} \int \tau \cdot dv [\text{cm}^{-2}], \quad (2)$$

where  $Q(T_{\text{ex}})$  is the partition function computed at the excitation temperature  $T_{\text{ex}}$ ,  $\nu$  is the frequency of the observed transition with the Einstein A-coefficient  $A_{ul}$  and the statistical weights of the lower  $g_1$  and upper levels  $g_u$ ;  $c$  is the speed of light, and  $k$  is the Boltzmann constant. The line opacity  $\tau$  was calculated from the measured brightness temperature  $T_{\text{mb}}$  and the temperature of the background continuum in a single side band  $T_c$ , using the relation  $\tau = -\ln\left(\frac{T_{\text{mb}}}{T_c}\right)$ .

$N_{\text{tot}}$  and  $N_1$  are the total column density and the column density in the lower state of transition. The  $N_1$  may be the same as total column density for the ground state lines when the excitation temperature is very low ( $T_{\text{ex}} \sim 2.73 \text{ K}$ ). Therefore we applied Eq. (2) to calculate column densities for the ground-state

<sup>5</sup> CASSIS (<http://cassis.cesr.fr>) has been developed by IRAP- UPS/CNRS..





**Fig. 2.** Average values of the line widths (*top panel*) and of the central velocity (*bottom panel*) from Gaussian fits for the observed emission lines of different molecules (“o” represents the outflow component). The emission lines of the envelope component are centred on  $-5.5 \text{ km s}^{-1}$ , as shown by the dashed line in the *bottom panel*.

transitions. For the absorptions that arise from the excited states we used Eq. (1) and assumed an excitation temperature of 10 K, as was derived for the foreground cloud in Paper I (see Table 4 in Van der Wiel et al. 2013).

The tentative absorption lines from a foreground cloud at  $V_{\text{lsr}} \sim 0 \text{ km s}^{-1}$  were observed of CCH (7<sub>7</sub>–6<sub>6</sub> at 611.265 GHz), CH (two transitions: 3/2<sub>2+</sub>–1/2<sub>1-</sub> at 532.724 and 3/2<sub>2-</sub>–1/2<sub>1+</sub> at 536.761 GHz), CH<sup>+</sup> (1–0 at 835.138 GHz), H<sub>2</sub>S (2<sub>12</sub>–1<sub>01</sub> at 736.034 GHz), NH<sub>3</sub> (1<sub>0</sub>–0<sub>0</sub> at 572.498 GHz), H<sub>2</sub>O (two transitions: 1<sub>10</sub>–1<sub>01</sub> at 556.936 and 1<sub>11</sub>–0<sub>00</sub> at 1113.343 GHz), OH<sup>+</sup> (three transitions:  $J = 2-1$ ,  $F = 3/2-1/2$  at 971.805,  $J = 1-1$ ,  $F = 3/2-1/2$  at 1033.004 and  $J = 1-1$ ,  $F = 3/2-3/2$  at 1033.119 GHz) and HF (1–0 at 1232.476 GHz). The estimated column densities for these species are listed in the upper part of Table 2.

Three broad absorptions are associated with the outflow (centred on  $\sim -13.8 \text{ km s}^{-1}$ ): H<sub>2</sub>O, CH<sup>+</sup> and HF. Their column densities are presented in the lower part of Table 2.

Bruderer et al. (2010b) analysed hydrides toward AFGL 2591 using HIFI. Our column density results agree well, with their measurements within the errors:  $3.1 \times 10^{13}$  and  $2.6 \times 10^{13} \text{ cm}^{-2}$  for CH,  $6.8 \times 10^{13}$  and  $1.8 \times 10^{14} \text{ cm}^{-2}$  for the CH<sup>+</sup> outflow component,  $1.1 \times 10^{14}$  and  $1.2 \times 10^{14} \text{ cm}^{-2}$  for CH<sup>+</sup>, and  $3.0 \times 10^{13}$  and  $6.1 \times 10^{13} \text{ cm}^{-2}$  for OH<sup>+</sup>, our results and from (Bruderer et al. 2010b) respectively. Bruderer et al. (2010b) also found lines of NH and H<sub>2</sub>O<sup>+</sup> in their spectra. These two species are not seen in our dataset, because of a slightly lower quality of spectral scans (Bruderer et al. 2010b have observations from the single frequency settings).

Based on *Herschel* data, Barlow et al. (2013) recently detected emission lines of <sup>36</sup>ArH<sup>+</sup> in the Crab nebula. Absorptions of this ion are also seen toward sources from *Herschel* Observations of EXtra-Ordinary Sources (HEXOS) and PRobing InterStellar Molecules with Absorption line Studies (PRISMAS) *Herschel* key programs (Schilke et al. 2014). The <sup>36</sup>ArH<sup>+</sup>  $J = 1-0$  transition at 617.525 GHz is not detected in our spectra. The upper limit of the column density is  $7.7 \times 10^{12} \text{ cm}^{-2}$  for the width of an absorption line of  $1 \text{ km s}^{-1}$ .

### 3.3. Emission line analysis

To estimate column densities and excitation temperatures from the observed emissions we constructed rotational diagrams assuming that all lines for a given molecule have the same excitation temperature. This method is a useful tool for estimating the column densities and the excitation temperatures when many transitions of a particular species are observed. However, in many cases its accuracy is limited because it is based on the assumption that the emission lines are optically thin and the emissions fill the beam.

Goldsmith & Langer (1999) improved this excitation analysis method by introducing correction factors for the effects of the beam dilution and optical depth. Using this *population diagram method*, we estimated the column density, the excitation temperature, and the emission extent for each molecule with the observed multiple transitions. With three free parameters (column density, excitation temperature, and the beam filling factor), we used this method only when at least four lines for a given molecule were observed. Otherwise, only the rotational method was applied. The rotation diagram gives beam-averaged column densities, while the population diagram gives source-averaged values. Hereafter, all stated column densities ( $N_{\text{col}}$ ) or excitation temperatures ( $T_{\text{ex}}$ ) were derived from the population diagrams, except those of HNC and N<sub>2</sub>H<sup>+</sup>, which were estimated from the rotational diagrams. At this point, the complementary JCMT data were crucial to increase the number of observed transitions for a given molecule.

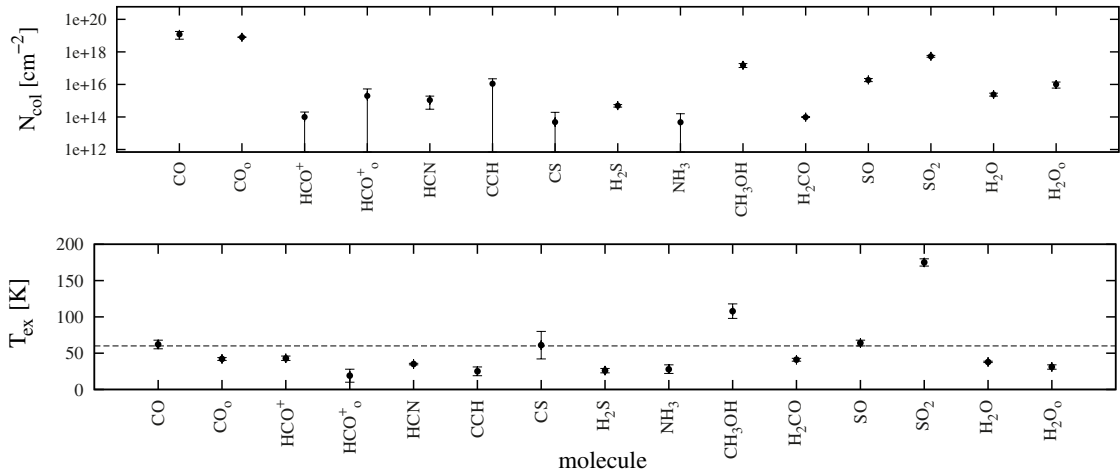
The column densities of CO, HCN and HCO<sup>+</sup> were obtained from their isotopologues (<sup>13</sup>CO, C<sup>18</sup>O, C<sup>17</sup>O, H<sup>13</sup>CN, HC<sup>15</sup>N, and H<sup>13</sup>CO<sup>+</sup>) using the standard isotopic ratios:  $^{12}\text{C}/^{13}\text{C} = 60$ ,  $^{16}\text{O}/^{18}\text{O} = 500$ ,  $^{16}\text{O}/^{17}\text{O} = 2500$ , and  $^{14}\text{N}/^{15}\text{N} = 270$  (Wilson & Rood 1994).

All column densities and excitation temperatures values based on the rotational and population diagrams methods are given in Table 3. The opacities and emission sizes for each molecule derived from the population diagrams are listed in Table 3 as well. It also contains information about the covered energy  $E_{\text{up}}$  range for a given species and the number of lines from different energy levels that were used for the analysis. The values of the excitation temperatures and column densities

**Table 3.** Parameters estimated from rotational and population diagrams methods (column densities, excitation temperatures, and emission extents) based on HIFI and JCMT data.

Molecule	$V$ [km s <sup>-1</sup> ]	$FWHM$ [km s <sup>-1</sup> ]	Rotational		Population			$\tau$	$E_{up}$ range [K]	No of trans.
			$N_{col}$ [cm <sup>-2</sup> ]	$T_{ex}$ [K]	$N_{col}$ [cm <sup>-2</sup> ]	$T_{ex}$ [K]	size[ $''$ ]			
CO	-4.8 $^{+0.3}_{-0.3}$	5.1 $^{+0.9}_{-0.9}$	6.0 $\times$ 10 <sup>16</sup> $^{+0.5}_{-0.5}$	162 $^{+10}_{-9}$	1.2 $\times$ 10 <sup>19</sup> $^{+0.4}_{-0.6}$	62 $^{+4}_{-6}$	17 $^{+3}_{-7}$	0.1–144	33–752	12
	-7.2 $^{+2.3}_{-2.3}$	15.4 $^{+1.8}_{-1.8}$	6.0 $\times$ 10 <sup>16</sup> $^{+1.0}_{-1.0}$	89 $^{+10}_{-9}$	8.0 $\times$ 10 <sup>18</sup> $^{+0.4}_{-0.4}$	42 $^{+2}_{-2}$	17 $^{+1}_{-1}$	0.01–34	33–752	12
HCO <sup>+</sup>	-5.6 $^{+0.2}_{-0.2}$	3.4 $^{+0.4}_{-0.4}$	2.3 $\times$ 10 <sup>13</sup> $^{+1.0}_{-1.0}$	35 $^{+3}_{-3}$	1.0 $\times$ 10 <sup>14</sup> $^{+1.0}_{-1.0}$	43 $^{+2}_{-3}$	11 $^{+1}_{-1}$	0.08–2.02	43–283	7
	-6.8 $^{+0.6}_{-0.6}$	7.3 $^{+2.0}_{-2.0}$	2.2 $\times$ 10 <sup>13</sup> $^{+4.2}_{-1.6}$	23 $^{+15}_{-15}$	2.0 $\times$ 10 <sup>15</sup> $^{+3.3}_{-1.6}$	19 $^{+9}_{-2}$	9.7 $^{+0.8}_{-2.4}$	0.7–35	43–154	4
HCN	-5.3 $^{+0.2}_{-0.2}$	4.2 $^{+0.7}_{-0.7}$	4.5 $\times$ 10 <sup>13</sup> $^{+0.7}_{-0.5}$	31 $^{+14}_{-10}$	1.1 $\times$ 10 <sup>15</sup> $^{+0.8}_{-0.7}$	35 $^{+1}_{-1}$	7.7 $^{+0.2}_{-0.2}$	0.2–4.4	43–234	6
HNC <sup>a</sup>	-5.4 $^{+0.1}_{-0.1}$	3.8 $^{+0.5}_{-0.5}$	4.8 $\times$ 10 <sup>12</sup> $^{+6.3}_{-0.5}$	43 $^{+9}_{-7}$					44–122	3
CCH	-6.3 $^{+0.5}_{-0.5}$	3.7 $^{+0.5}_{-0.5}$	2.2 $\times$ 10 <sup>14</sup> $^{+0.3}_{-0.2}$	22 $^{+3}_{-3}$	1.1 $\times$ 10 <sup>16</sup> $^{+1.1}_{-0.5}$	25 $^{+6}_{-5}$	5.3 $^{+0.8}_{-0.6}$	0.3–7.1	42–151	4
CN <sup>a</sup>	-5.6 $^{+0.2}_{-0.2}$	3.2 $^{+0.5}_{-0.5}$	9.7 $\times$ 10 <sup>13</sup> $^{+0.8}_{-0.7}$	22 $^{+1}_{-1}$	1.3 $\times$ 10 <sup>14</sup> $^{+0.3}_{-0.3}$	26 $^{+4}_{-3}$	23 $^{+6}_{-4}$	0.01–0.18	33–114	3
CS	-5.5 $^{+0.4}_{-0.4}$	3.9 $^{+0.5}_{-0.5}$	7.4 $\times$ 10 <sup>13</sup> $^{+0.5}_{-0.4}$	26 $^{+12}_{-10}$	4.9 $\times$ 10 <sup>13</sup> $^{+14.6}_{-0.6}$	61 $^{+7}_{-19}$	14 $^{+4}_{-9}$	0.01–0.09	66–282	7
H <sub>2</sub> S	-5.7 $^{+0.7}_{-0.7}$	3.4 $^{+0.8}_{-0.8}$	1.1 $\times$ 10 <sup>13</sup> $^{+1.1}_{-0.7}$	56 $^{+25}_{-14}$	4.9 $\times$ 10 <sup>14</sup> $^{+0.9}_{-0.6}$	26 $^{+3}_{-2}$	8.9 $^{+0.6}_{-0.7}$	0.01–5.6	55–350	5
NH <sub>3</sub>	-5.1 $^{+0.6}_{-0.6}$	4.1 $^{+1.0}_{-1.0}$	2.8 $\times$ 10 <sup>13</sup> $^{+0.3}_{-0.2}$	67 $^{+6}_{-5}$	4.8 $\times$ 10 <sup>13</sup> $^{+11}_{-2.4}$	28 $^{+3}_{-6}$	9.6 $^{+1.8}_{-0.6}$	0.1–1.6	28–170	5
N <sub>2</sub> H <sup>+</sup> <sup>a</sup>	-5.9 $^{+0.2}_{-0.2}$	2.8 $^{+0.3}_{-0.3}$	5.6 $\times$ 10 <sup>11</sup> $^{+0.1}_{-1.6}$	19 $^{+13}_{-3}$					45–125	3
NO <sup>a</sup>	-4.8 $^{+0.5}_{-0.5}$	5.8 $^{+2.8}_{-2.8}$	7.2 $\times$ 10 <sup>15</sup> $^{+0.7}_{-0.5}$	25 $^{+14}_{-10}$	1.7 $\times$ 10 <sup>16</sup> $^{+3.6}_{-0.5}$	54 $^{+9}_{-14}$	12 $^{+15}_{-6}$	0.015–0.021	36–115	2
CH <sub>3</sub> OH	-5.7 $^{+0.5}_{-0.5}$	3.3 $^{+0.6}_{-0.6}$	1.8 $\times$ 10 <sup>14</sup> $^{+1.0}_{-0.7}$	209 $^{+171}_{-62}$	1.5 $\times$ 10 <sup>17</sup> $^{+0.4}_{-0.3}$	108 $^{+10}_{-7}$	1.5 $^{+0.1}_{-0.1}$	0.6–10.4	25–352	49
H <sub>2</sub> CO	-5.4 $^{+0.3}_{-0.3}$	3.6 $^{+0.7}_{-0.7}$	2.0 $\times$ 10 <sup>13</sup> $^{+1.0}_{-0.9}$	34 $^{+12}_{-6}$	9.9 $\times$ 10 <sup>13</sup> $^{+0.1}_{-0.1}$	41 $^{+2}_{-2}$	7.3 $^{+0.1}_{-0.1}$	0.02–0.61	32–263	14
SO	-5.5 $^{+0.4}_{-0.4}$	4.9 $^{+0.7}_{-0.7}$	1.5 $\times$ 10 <sup>14</sup> $^{+0.2}_{-0.2}$	53 $^{+23}_{-20}$	1.9 $\times$ 10 <sup>16</sup> $^{+0.4}_{-0.3}$	64 $^{+2}_{-4}$	2.7 $^{+0.3}_{-0.2}$	0.1–6.1	26–405	22
SO <sub>2</sub>	-5.1 $^{+0.4}_{-0.4}$	4.6 $^{+0.9}_{-0.9}$	3.0 $\times$ 10 <sup>14</sup> $^{+2.0}_{-1.7}$	92 $^{+77}_{-43}$	5.4 $\times$ 10 <sup>17</sup> $^{+0.7}_{-0.6}$	175 $^{+5}_{-4}$	0.9 $^{+0.1}_{-0.1}$	0.5–8.9	31–354	47
H <sub>2</sub> O	-4.8 $^{+0.9}_{-0.9}$	3.1 $^{+0.6}_{-0.6}$	3.5 $\times$ 10 <sup>13</sup> $^{+3.0}_{-1.4}$	63 $^{+17}_{-15}$	2.4 $\times$ 10 <sup>15</sup> $^{+0.3}_{-0.5}$	38 $^{+1}_{-1}$	9.1 $^{+0.5}_{-0.6}$	0.4–104	53–305	8
	-6.0 $^{+0.2}_{-0.2}$	12.1 $^{+2.3}_{-2.3}$	5.5 $\times$ 10 <sup>13</sup> $^{+3.1}_{-1.9}$	43 $^{+35}_{-20}$	1.0 $\times$ 10 <sup>16</sup> $^{+0.4}_{-0.4}$	31 $^{+3}_{-3}$	4.9 $^{+0.3}_{-0.4}$	0.2–5.7	101–305	6

**Notes.** The second column shows the values of the central velocity of the observed lines. When there are two values, the first corresponds to the envelope component, the second to the outflow. The last 3 columns show first the range of optical depth  $\tau$  for observed lines, then the  $E_{up}$  range, which is covered by observed features, and finally the number of lines from different energy levels used for the analysis; i.e., we observed 4 lines of NO, but they originate only from 2 different energy levels. Population diagram method was used when at least 4 lines of a given molecule were observed, which is why there are no values for HNC and N<sub>2</sub>H<sup>+</sup>. <sup>(a)</sup> Indicates higher uncertainty of measurements because e.g. only 3 different levels were observed.

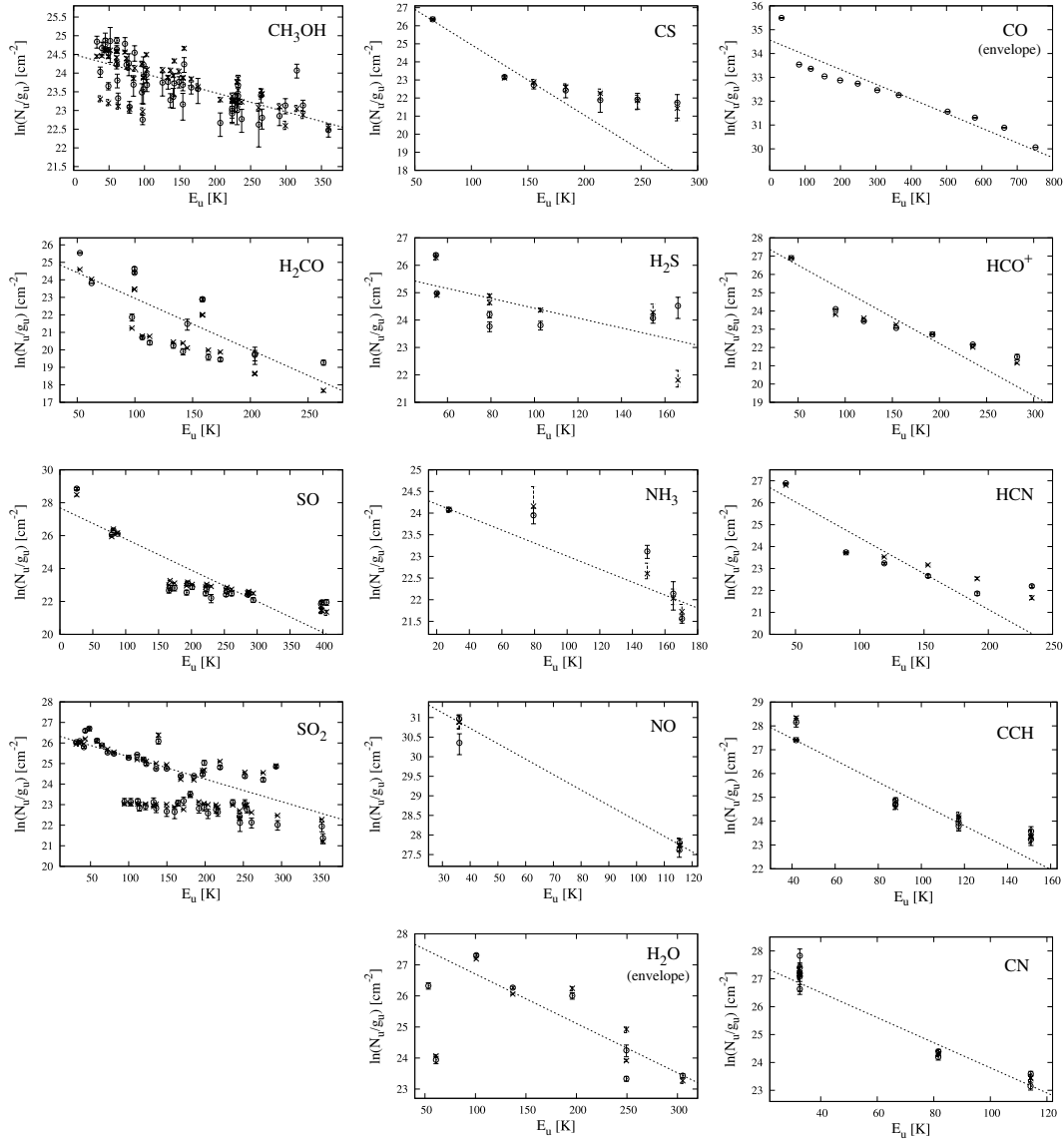
**Fig. 3.** Column densities and excitation temperatures estimated from the population diagrams, without the uncertain measurements, “o” represents the outflow component.

are plotted in Fig. 3, excluding the uncertain measurements (i.e. HNC, N<sub>2</sub>H<sup>+</sup>, CN, and NO).

Based on the optical depths values from Table 3, the lines of CN, CS, NO, and H<sub>2</sub>CO can be characterised as optically thin ( $\tau < 0.6$ ). However, the results of CN and NO are uncertain because of only a few lines were observed. For optically thin lines calculations based on the rotational diagrams resulted in

good approximations of the column densities and the excitation temperatures. The other molecular lines were characterised as optically thick. For these molecular species the population diagram method was more accurate.

The emission extent of the analysed molecules associated with AFGL 2591 ranges from around 2 $''$  (species such as SO,



**Fig. 4.** Population diagrams. Open circles represent the observational data and crosses are the best-fit model from population diagram analysis. Dotted lines correspond to a linear fit to the rotational diagram.

SO<sub>2</sub>, and CH<sub>3</sub>OH) up to 23'' (CN). For most species emission sizes are smaller than 17''.

From the comparison of the temperatures derived from the population diagrams (see the bottom panel of Fig. 3) it is possible to distinguish warm (e.g. CH<sub>3</sub>OH and SO<sub>2</sub>) and cold (e.g. HCN, H<sub>2</sub>S, and NH<sub>3</sub>) species. As cold species we classify those with excitation temperatures of up to 70 K. Warm molecules have higher temperatures, up to 175 K for SO<sub>2</sub>. It is difficult to give an accurate borderline here and classify all species, but the wide range of excitation temperatures seems significant. Moreover, it was shown before by Bisschop et al. (2007) that some of the complex organic species can be classified as both warm and cold, which may indicate that they are present in multiple physical components.

The population diagrams are presented in Fig. 4. They show evidence for an excitation gradient of several species (HCO<sup>+</sup>, HCN, CS, and SO), which means that the population diagram method may be not enough to analyse all observed molecules. This is a motivation to use more sophisticated method in the near future (i.e., radiative transfer modelling) to study our spectral survey.

## 4. Discussion

### 4.1. CI and CII

C and C<sup>+</sup> are the only atomic species found in our HIFI spectral survey of AFGL 2591. Both fine-structure transitions of neutral carbon, <sup>3</sup>P<sub>1</sub>–<sup>3</sup>P<sub>0</sub> at 492 GHz and <sup>3</sup>P<sub>2</sub>–<sup>3</sup>P<sub>1</sub> at 809 GHz, were observed towards AFGL 2591. These transitions consist of two components originating from the envelope and the outflow, similar to the CO lines (see Fig. A.1). CI was observed previously in AFGL 2591 by Van der Tak et al. (1999), but [CII] was observed for the first time with *Herschel*. The [CII] <sup>2</sup>P<sub>3/2</sub>–<sup>2</sup>P<sub>1/2</sub> line, an important interstellar coolant, shows several velocity components, two of them correspond to those in CI and CO. The [CII] line profile is distorted by a contamination from the off-position even after applying corrections within HIPE (Fig. A.1).

### 4.2. CO and its isotopologues

CO is one of the most often studied molecules (e.g. Mitchell et al. 1989; Black et al. 1990; Hasegawa & Mitchell 1995).

Based on CO observation, Lada et al. (1984) found an extended bipolar outflow associated with AFGL 2591. Many strong lines of CO and its isotopologues ( $^{13}\text{CO}$ ,  $\text{C}^{18}\text{O}$ ,  $\text{C}^{17}\text{O}$ ) were also detected in our HIFI spectra, clearly showing the envelope and outflow components. The  $\text{C}^{17}\text{O}$  lines are weaker and show only the envelope components. The abundance of CO =  $3 \times 10^{-5}$  was calculated in Paper I. The CO column density in this work was estimated at  $1.2 \times 10^{19} \text{ cm}^{-2}$ ; Van der Tak et al. (2000b) derived a similar value of  $3.4 \times 10^{19} \text{ cm}^{-2}$ .

#### 4.3. $\text{HCO}^+$

$\text{HCO}^+$  was identified by intense lines in the HIFI and JCMT spectra. Moreover, three lines of  $\text{H}^{13}\text{CO}^+$  were also positively detected. The abundance of  $\text{HCO}^+$  was estimated at  $9 \times 10^{-9}$  (Paper I) and column density at  $1.0 \times 10^{14} \text{ cm}^{-2}$ . Carr et al. (1995) estimated an abundance of  $4 \times 10^{-10}$  and Van der Tak et al. (1999) derived  $[\text{HCO}^+] = 1 \times 10^{-8}$  by using a model with lower  $\text{H}_2$  column density.

#### 4.4. N-bearing species

Six N-bearing species were observed in the HIFI spectra: HCN, HNC, CN, NO,  $\text{N}_2\text{H}^+$ , and  $\text{NH}_3$ . All of these molecules have been detected before in AFGL 2591 (e.g. Takano et al. 1986; Carr et al. 1995; Boonman et al. 2001). Lines of N-bearing species observed with the *Herschel*/HIFI are weaker than those of CO and were fitted with a single-Gaussian profile, revealing these species to be components of the protostellar envelope, centred on  $-5.5 \text{ km s}^{-1}$ . Only o- $\text{NH}_3$  shows a tentative absorption feature from a foreground cloud at  $V_{\text{lsr}} = 0 \text{ km s}^{-1}$ . Two features observed with the JCMT, HCN 4–3, and HNC 4–3 show a contribution from the outflow and double-Gaussian profiles were fitted to these lines. We did not find NH and  $\text{NH}_2$ , which were detected in other HIFI spectral surveys (e.g. Zernickel et al. 2012). Upper limits are  $0.8 \text{ K km s}^{-1}$  for the NH 1–0 line near 946 GHz and  $0.6 \text{ K km s}^{-1}$  for the  $\text{NH}_2$  1–0 line near 953 GHz. Upper limits were measured in the same way as in Paper I, that is, considering  $3 \text{ km s}^{-1}$  a typical line width, hence using  $5\sigma_{\text{rms}} \times 3 \text{ km s}^{-1}$ . Among the observed features, two lines of vibrationally excited HCN 4–3,  $\nu = 1c$  and  $\nu = 1d$  are found (JCMT data). Line  $\nu = 1c$  was observed before by Van der Tak et al. (1999). Boonman et al. (2001) analysed excited HCN, the 4–3 and 9–8 transitions. The interferometric observations from Veach et al. (2013) showed vibrationally excited  $\nu = 1$  and also  $\nu = 2$  HCN 4–3 lines. These authors suggested that the  $\nu = 2$  HCN lines might be a useful tool to study a protostellar disk. Takano et al. (1986) observed ammonia transitions (1, 1) and (2, 2) with the Effelsberg 100 m telescope. They found a compact  $\text{NH}_3$  cloud of around 0.6 pc diameter around the central source. These authors estimated a column density of  $8 \times 10^{13} \text{ cm}^{-2}$ . In comparison, calculations of our work gave a column density of  $4.8 \times 10^{13} \text{ cm}^{-2}$ .

#### 4.5. S-bearing species

Of the S-bearing molecules we detected with HIFI CS,  $\text{H}_2\text{S}$ ,  $\text{H}_2^{34}\text{S}$ , SO, and  $\text{SO}_2$ . All of these molecules have been observed before in AFGL 2591 (e.g. Yamashita et al. 1987; Van der Tak et al. 2003; Bruderer et al. 2009). Additionally, from the JCMT dataset we have several lines of these molecules, and also isotopologues of CS, SO, and  $\text{SO}_2$  ( $^{13}\text{CS}$  and  $\text{C}^{34}\text{S}$ ,  $^{34}\text{SO}$ ,  $^{34}\text{SO}_2$ ), as well as OCS and o- $\text{H}_2\text{CS}$ . SO and  $\text{SO}_2$  show many weak lines of the envelope component.  $\text{SO}_2$  is an example of warm species with an excitation temperature of 175 K, whereas

$\text{H}_2\text{S}$  is classified as colder species with an excitation temperature of 26 K. CS and SO have similar excitation temperatures, 61 K and 64 K. Van der Tak et al. (2003) studied the sulphur chemistry in the envelopes of massive star forming regions and found excitation temperatures of 185 K for  $\text{SO}_2$ , which is a similar result to the one calculated in this work. However, the column density of  $\text{SO}_2$  varies strongly,  $5.2 \times 10^{14} \text{ cm}^{-2}$  Van der Tak et al. (2003),  $5.4 \times 10^{17} \text{ cm}^{-2}$  our work. Results of column density of CS also differ in one order of magnitude,  $3 \times 10^{13} \text{ cm}^{-2}$  and  $4.9 \times 10^{14} \text{ cm}^{-2}$  (Van der Tak et al. 2003) and our work, respectively. The population diagram method is a good first step for the spectral surveys analysis, but in some cases a more advanced method is needed. Especially when there are not enough observed transitions from the lower energy levels for a given molecule, for example SO or CS and the excitation gradient is visible (see Fig. 4). We are planning to use radiative transfer modelling and estimate molecular abundances in the near future.

#### 4.6. CCH, CH, $\text{CH}^+$ , OH, and $\text{OH}^+$

Our spectra also revealed lines from the protostellar envelope and foreground clouds that belong to CCH, CH,  $\text{CH}^+$ , OH, and  $\text{OH}^+$ . CCH and CH show three absorption lines at  $\sim 0 \text{ km s}^{-1}$ ,  $\text{OH}^+$  three absorptions at  $\sim 3.6 \text{ km s}^{-1}$ . Using HIFI, Bruderer et al. (2010b,a) found lines of CH,  $\text{CH}^+$ , NH,  $\text{OH}^+$ , and  $\text{H}_2\text{O}^+$ , while lines of  $\text{NH}^+$  and  $\text{SH}^+$  have not been detected. Bruderer et al. (2010b) concluded that absorption lines of NH,  $\text{OH}^+$  and  $\text{H}_2\text{O}^+$  originate from a foreground cloud and an outflow lobe, while the emission lines of CH and  $\text{CH}^+$  are connected with the protostellar envelope (compare Sect. 3.2).

#### 4.7. Water

Water lines have also been detected in our spectra. We found four transitions of o- $\text{H}_2\text{O}$  ( $1_{10}-1_{01}$  at 557 GHz,  $3_{12}-3_{03}$  at 1097 GHz,  $3_{12}-2_{21}$  at 1153 GHz, and  $3_{21}-3_{12}$  at 1163 GHz) and four transitions of p- $\text{H}_2\text{O}$  ( $2_{11}-2_{02}$  at 752 GHz,  $2_{02}-1_{11}$  at 988 GHz,  $1_{11}-0_{00}$  at 1113 GHz, and  $2_{20}-2_{11}$  at 1229 GHz). They show different profiles, mostly the envelope and outflow components, but also some absorptions (see Fig. A.1). For the envelope component we estimated a column density of  $2.4 \times 10^{15} \text{ cm}^{-2}$ , an excitation temperature of 38 K, and an emission extent of  $9.1''$ . The full analysis of water lines in AFGL 2591 as part of the Water In Star forming regions with *Herschel* (WISH) Project will be presented in the forthcoming paper of Choi et al. (2014).

#### 4.8. HF

HF is the only detected fluorine-bearing species in AFGL 2591. Its 1–0 transition at 1233 GHz was observed and analysed by Emprechtinger et al. (2012). They calculated an HF column density of  $2 \times 10^{14} \text{ cm}^{-2}$  and  $4 \times 10^{13} \text{ cm}^{-2}$  for emission and absorption.

#### 4.9. HCl

Thanks to HIFI, many chlorine-bearing molecules (e.g. HCl,  $\text{H}^{37}\text{Cl}$ ,  $\text{H}_2\text{Cl}^+$ ,  $\text{H}_2^{37}\text{Cl}^+$ ) were observed in different environments, for example toward protostellar shocks (Codella et al. 2012), diffuse clouds (Monje et al. 2013), and star forming regions (Neufeld et al. 2012). HCl and  $\text{H}^{37}\text{Cl}$  are the only observed chlorine-bearing species in our HIFI spectra of AFGL 2591. Three hyperfine components of HCl from the energy level of  $E_{\text{up}} = 30 \text{ K}$  and two from the higher state  $E_{\text{up}} = 90.1 \text{ K}$  were detected. In agreement with



Neufeld et al. (2012), neither lines of  $\text{H}_2\text{Cl}^+$  nor lines of  $\text{H}_2^{37}\text{Cl}^+$  toward AFGL 2591 were found.

#### 4.10. Complex species

From the HIFI spectral survey we found only two molecules (methanol and formaldehyde) that belong to complex organics. Bisschop et al. (2007) showed before that AFGL 2591 is a line-poor source. These authors analysed complex organic molecules in massive young stellar objects and found only a few of them in AFGL 2591; all of the intensities of the observed lines were very low. Many weak  $\text{CH}_3\text{OH}$  and  $\text{H}_2\text{CO}$  lines were detected in our HIFI spectra. Their column densities and excitation temperatures are  $1.5 \times 10^{17} \text{ cm}^{-2}$  and 108 K for  $\text{CH}_3\text{OH}$ , and  $9.9 \times 10^{13} \text{ cm}^{-2}$  and 41 K for  $\text{H}_2\text{CO}$ . Van der Tak et al. (2000a) estimated  $1.2 \times 10^{15} \text{ cm}^{-2}$  and 163 K for  $\text{CH}_3\text{OH}$ , and  $8.0 \times 10^{13} \text{ cm}^{-2}$  and 89 K for  $\text{H}_2\text{CO}$ . From the rotational diagrams Bisschop et al. (2007) derived  $4.7 \times 10^{16} \text{ cm}^{-2}$  and 147 K for methanol. All of these results vary slightly, but also suggest that methanol represents warm species.

## 5. Conclusions

The main conclusions from our AFGL 2591 spectral survey are as follows:

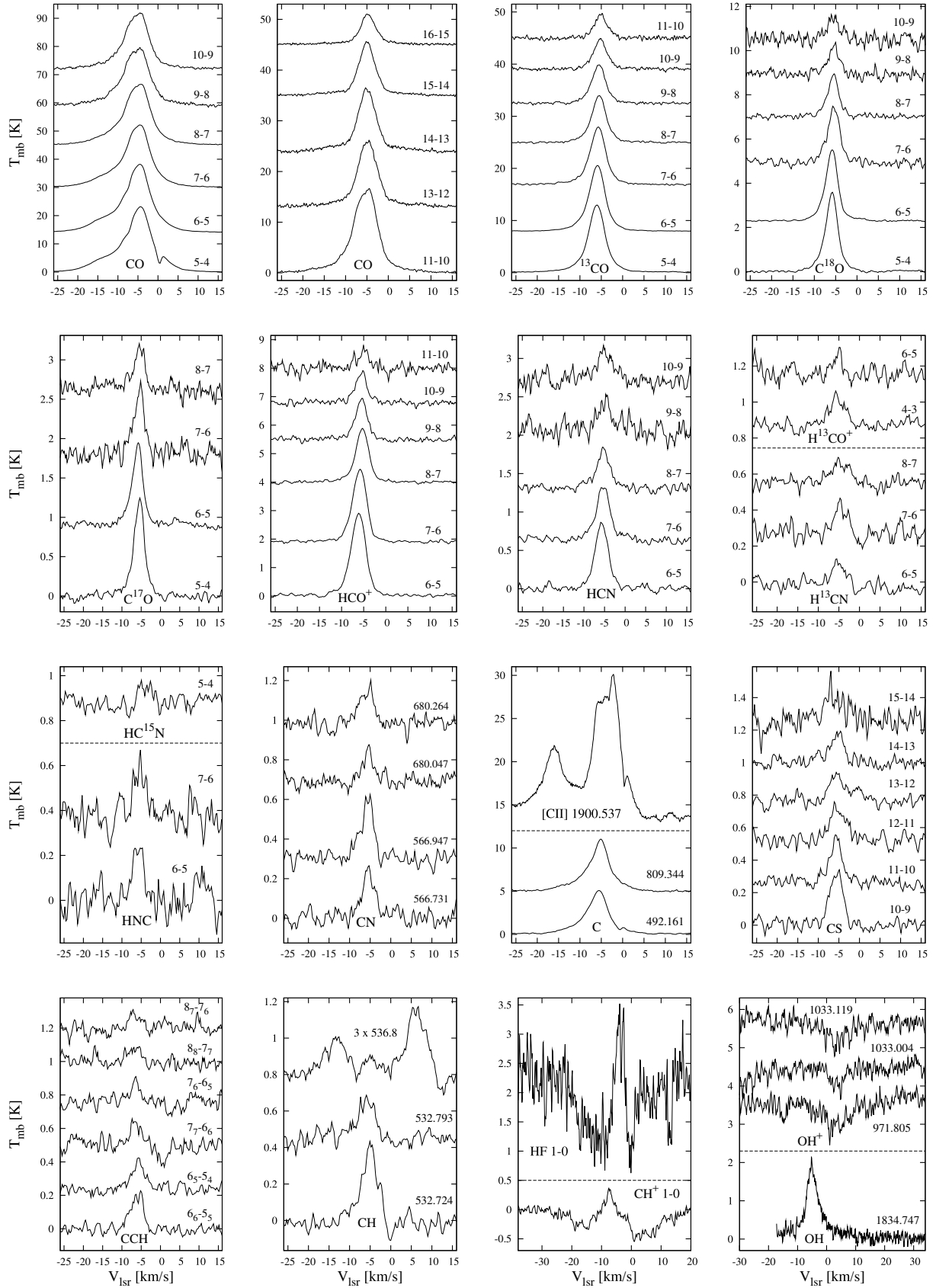
1. In the *Herschel*/HIFI spectral survey of AFGL 2591 we observed 268 lines (excluding blends) of a total of 32 species. No unidentified features were found in the spectra. JCMT data supplemented the excitation analysis of several species seen in emissions.
2. Among the observed 268 lines, 16 absorptions were detected. Most of them belong to the known foreground cloud at  $V_{\text{lsr}} \sim 0 \text{ km s}^{-1}$ . Three broad absorptions are associated with the outflow lobe. The estimated column densities agree well with previous work.
3. Based on the population diagram method, the column densities and excitation temperatures were estimated. Molecular column densities range from  $6 \times 10^{11}$  to  $1 \times 10^{19} \text{ cm}^{-2}$  and excitation temperatures range from 19 to 175 K. We can distinguish between species of higher (e.g.  $\text{CH}_3\text{OH}$ , and  $\text{SO}_2$ ) and lower (e.g.  $\text{HCN}$ ,  $\text{H}_2\text{S}$ , and  $\text{NH}_3$ ) excitation temperature.
4. The population diagram method is a very useful tool for spectral survey analysis, but it is far from being perfect. Several species ( $\text{HCO}^+$ ,  $\text{HCN}$ ,  $\text{CS}$ , and  $\text{SO}$ ) show evidence of an excitation gradient, which is a motivation to use a more sophisticated method (i.e., radiative transfer modelling) in the near future to study molecules observed in the protostellar envelope of AFGL 2591.

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## Appendix A: HIFI/CHESS spectral survey



**Fig. A.1.** Line profiles of all identified species in HIFI spectral survey of AFGL 2591. Molecules (e.g. HCl, CH) for which hyperfine components were detected are plotted all together and are centred on the velocity of the middle line. Lines with multiply profiles (e.g. CO – envelope and outflow components; HF – emission and absorptions) are also presented in one figure and are centered at the velocity of the envelope component; please compare with Table A.1.

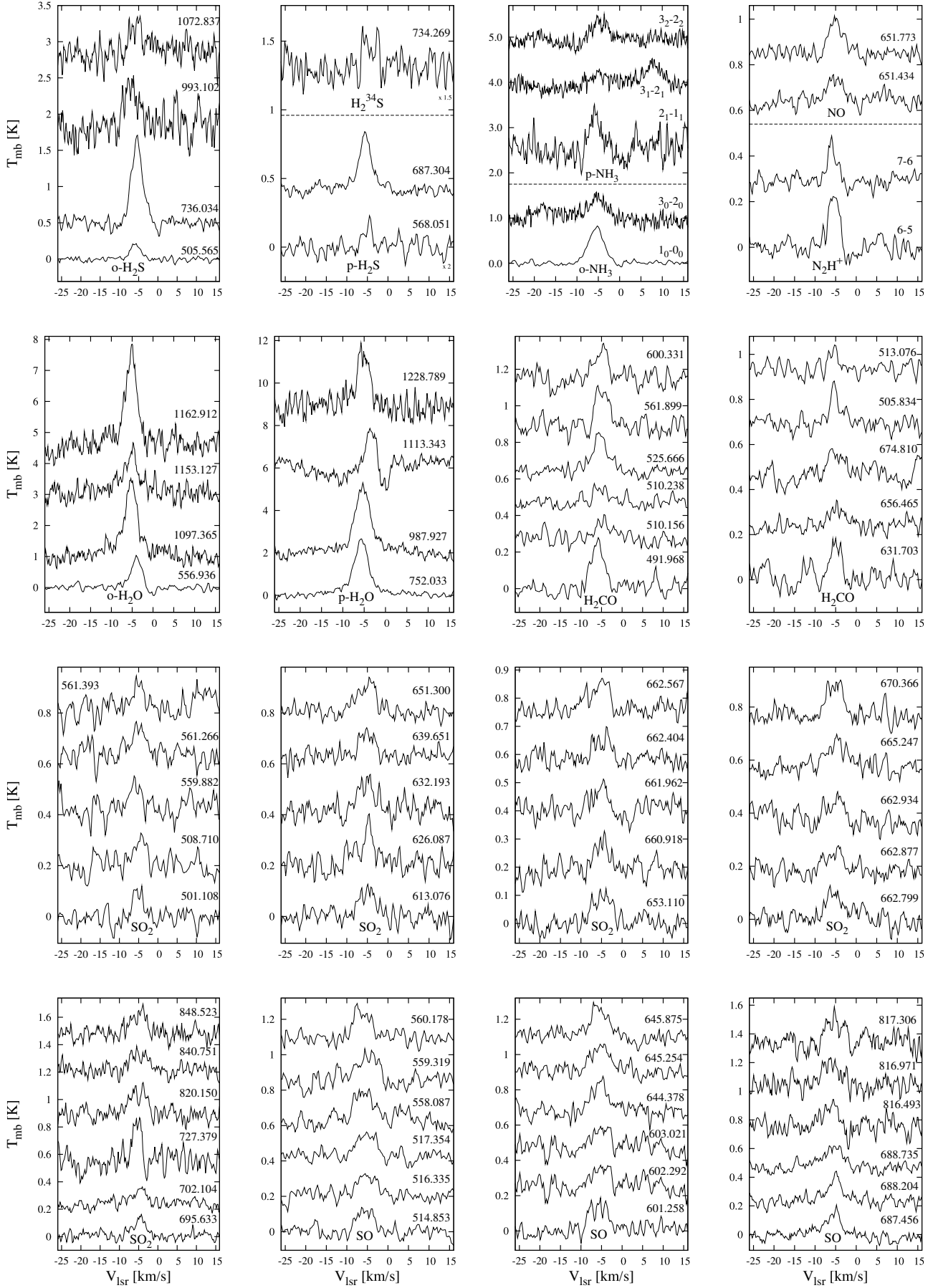


Fig. A.1. continued.

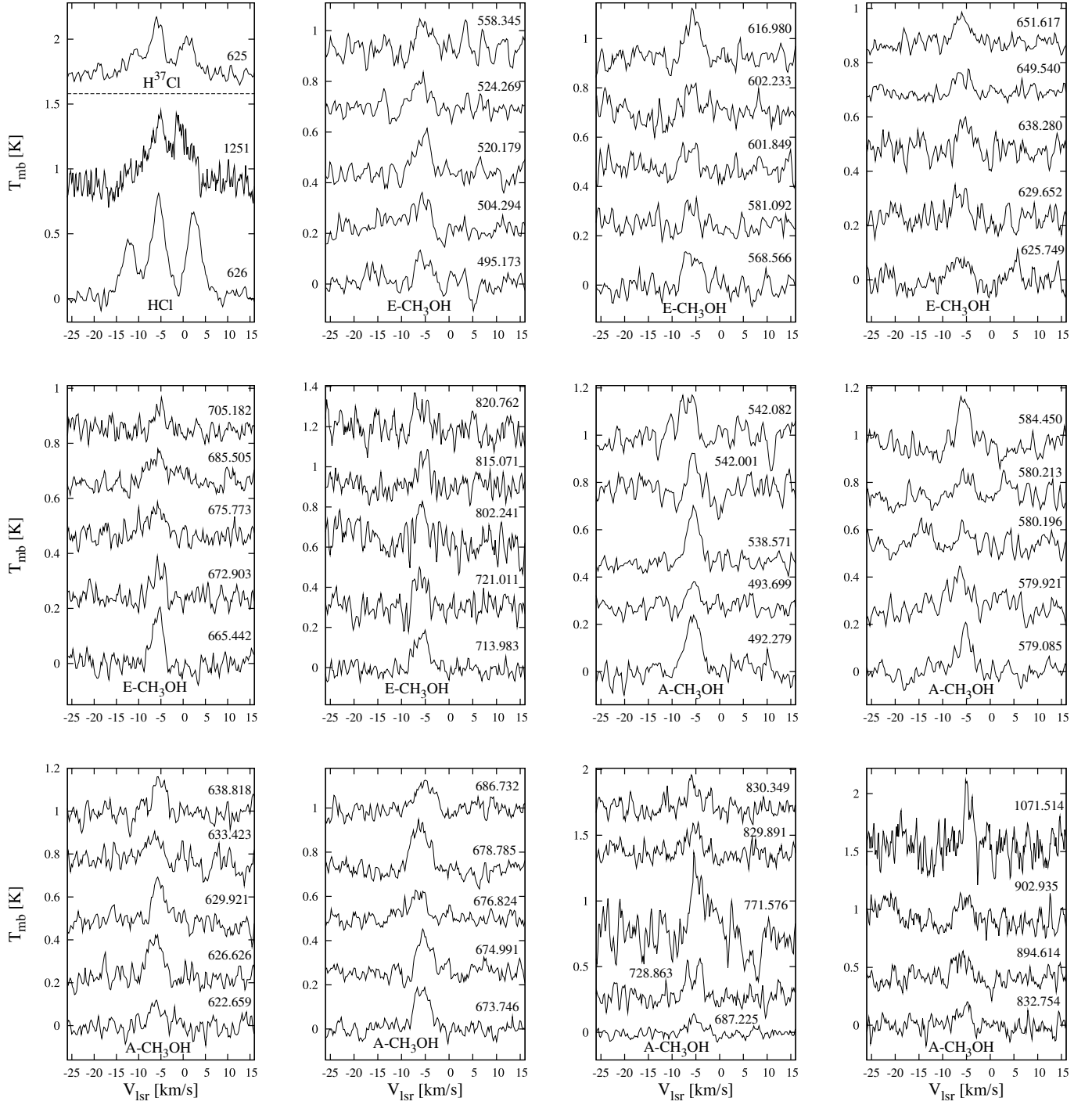


Fig. A.1. continued.



**Table A.1.** Results of line measurements of all identified species from the HIFI spectral survey of AFGL 2591 ( \* indicates JCMT data).

Transition	Frequency [GHz]	$E_{\text{up}}/k$ [K]	$V_{\text{lsr}}$ [km s <sup>-1</sup> ]	$\Delta V$ [km s <sup>-1</sup> ]	$\int T_{\text{mb}} dV$ [K km s <sup>-1</sup> ]	$T_{\text{peak}}$ [K]
<b>CO</b>						
3–2*	345.796	33.2	–4.16 ± 0.01 –14.03 ± 0.01	4.13 ± 0.01 11.40 ± 0.02	195.3 ± 0.2 170.8 ± 0.4	44.40 ± 0.02 14.08 ± 0.01
5–4	576.268	83.0	–4.42 ± 0.02 –6.70 ± 0.06	4.98 ± 0.05 15.85 ± 0.12	76.6 ± 0.1 155.7 ± 0.2	14.45 ± 0.14 9.23 ± 0.13
6–5	691.473	116.2	–4.61 ± 0.01 –7.23 ± 0.05	5.39 ± 0.04 15.70 ± 0.10	92.1 ± 0.1 145.5 ± 0.1	16.07 ± 0.12 8.71 ± 0.11
7–6	806.652	154.9	–4.78 ± 0.01 –7.36 ± 0.04	5.52 ± 0.03 15.09 ± 0.09	90.8 ± 0.1 119.5 ± 0.1	15.45 ± 0.09 7.44 ± 0.09
8–7	921.800	199.1	–4.90 ± 0.02 –7.21 ± 0.09	5.96 ± 0.06 15.17 ± 0.18	100.1 ± 0.2 105.9 ± 0.2	15.77 ± 0.17 6.56 ± 0.17
9–8	1036.912	248.9	–5.02 ± 0.03 –7.22 ± 0.18	6.43 ± 0.09 15.96 ± 0.42	109.9 ± 0.3 77.8 ± 0.5	16.06 ± 0.27 4.58 ± 0.27
10–9	1151.985	304.2	–4.97 ± 0.02 –7.29 ± 0.17	6.01 ± 0.07 14.64 ± 0.35	102.9 ± 0.2 66.1 ± 0.4	16.08 ± 0.23 4.24 ± 0.23
11–10	1267.014	365.0	–5.05 ± 0.02 –7.21 ± 0.17	5.71 ± 0.06 14.87 ± 0.40	100.8 ± 0.2 58.1 ± 0.4	10.10 ± 0.12 2.23 ± 0.12
13–12	1496.923	503.1	–4.94 ± 0.01 –4.95 ± 0.30	4.82 ± 0.03 18.24 ± 0.84	69.5 ± 0.1 34.7 ± 0.8	9.34 ± 0.05 1.23 ± 0.06
14–13	1611.794	580.5	–5.00 ± 0.01 –5.98 ± 0.11	4.54 ± 0.02 17.47 ± 0.36	62.2 ± 0.1 24.7 ± 0.4	8.80 ± 0.03 0.91 ± 0.03
15–14	1726.603	663.4	–5.08 ± 0.01 –6.07 ± 0.13	4.22 ± 0.02 16.61 ± 0.41	46.6 ± 0.1 18.0 ± 0.4	7.01 ± 0.03 0.69 ± 0.03
16–15	1841.346	751.7	–4.82 ± 0.01 –4.96 ± 0.11	3.80 ± 0.03 13.34 ± 0.40	23.1 ± 0.1 12.1 ± 0.4	3.83 ± 0.03 0.57 ± 0.03
<b><sup>13</sup>CO</b>						
3–2*	330.588	31.3	–5.58 ± 0.01 –7.33 ± 0.01	3.68 ± 0.02 9.13 ± 0.04	92.3 ± 0.6 97.3 ± 1.1	23.56 ± 0.05 10.01 ± 0.02
5–4	550.926	79.3	–5.85 ± 0.01 –6.34 ± 0.04	3.78 ± 0.03 9.06 ± 0.14	38.7 ± 0.1 31.6 ± 0.2	9.61 ± 0.12 3.27 ± 0.12
6–5	661.067	111.1	–5.76 ± 0.01 –6.41 ± 0.03	3.58 ± 0.03 8.21 ± 0.11	34.6 ± 0.1 31.0 ± 0.2	9.07 ± 0.11 3.54 ± 0.11
7–6	771.184	148.1	–5.66 ± 0.01 –6.34 ± 0.06	3.63 ± 0.04 8.06 ± 0.19	32.1 ± 0.2 22.4 ± 0.2	8.31 ± 0.15 2.61 ± 0.16
8–7	881.273	190.4	–5.55 ± 0.01 –5.92 ± 0.04	3.18 ± 0.05 7.03 ± 0.17	21.0 ± 0.2 20.8 ± 0.2	6.21 ± 0.16 2.78 ± 0.17
9–8	991.329	237.9	–5.43 ± 0.02 –5.77 ± 0.05	2.94 ± 0.10 6.03 ± 0.21	14.1 ± 0.3 18.5 ± 0.4	4.51 ± 0.28 2.88 ± 0.29
10–9	1101.350	290.8	–5.33 ± 0.04 –5.41 ± 0.04	2.32 ± 0.18 5.19 ± 0.21	6.2 ± 0.4 18.1 ± 0.4	2.53 ± 0.32 3.28 ± 0.33
11–10	1211.330	348.9	–5.22 ± 0.03	3.81 ± 0.07	17.4 ± 0.1	4.29 ± 0.07
<b>C<sup>18</sup>O</b>						
5–4	548.831	79.0	–5.79 ± 0.01 –6.30 ± 0.04	2.55 ± 0.05 5.72 ± 0.12	6.2 ± 0.1 7.8 ± 0.1	2.30 ± 0.07 1.29 ± 0.07
6–5	658.553	110.6	–5.69 ± 0.01 –6.23 ± 0.06	2.63 ± 0.06 5.84 ± 0.19	6.3 ± 0.1 6.2 ± 0.2	2.26 ± 0.08 0.99 ± 0.08
7–6	768.252	147.5	–5.47 ± 0.05 –6.01 ± 0.17	2.58 ± 0.19 5.71 ± 0.51	4.6 ± 0.3 5.3 ± 0.5	1.69 ± 0.19 0.87 ± 0.20
8–7	877.922	189.6	–5.41 ± 0.04 –5.82 ± 0.06	1.79 ± 0.15 4.32 ± 0.18	1.7 ± 0.2 4.9 ± 0.2	0.88 ± 0.10 1.07 ± 0.10
9–8	987.560	237.0	–5.30 ± 0.03	3.24 ± 0.06	4.1 ± 0.1	1.19 ± 0.07
10–9	1097.163	289.7	–5.51 ± 0.08	4.00 ± 0.19	3.8 ± 0.2	0.90 ± 0.05
<b>C<sup>17</sup>O</b>						
3–2*	337.061	32.4	–5.64 ± 0.04 –6.90 ± 0.20	2.80 ± 0.10 5.70 ± 0.30	9.8 ± 0.6 7.3 ± 1.0	3.29 ± 0.08 1.20 ± 0.04
5–4	561.713	80.9	–5.63 ± 0.03	3.44 ± 0.07	4.3 ± 0.1	1.17 ± 0.02
6–5	674.009	113.2	–5.81 ± 0.03	3.51 ± 0.08	3.6 ± 0.1	0.97 ± 0.02
7–6	786.281	150.9	–5.60 ± 0.07	3.37 ± 0.16	2.8 ± 0.2	0.79 ± 0.03
8–7	898.523	194.1	–5.44 ± 0.06	3.04 ± 0.15	1.7 ± 0.1	0.51 ± 0.02
<b>C</b>						
<sup>3</sup> P <sub>1</sub> – <sup>3</sup> P <sub>0</sub>	492.161	23.6	–5.72 ± 0.02 –7.17 ± 0.06	3.97 ± 0.06 10.19 ± 0.15	13.66 ± 0.39 19.87 ± 0.79	3.23 ± 0.05 1.83 ± 0.05
<sup>3</sup> P <sub>2</sub> – <sup>3</sup> P <sub>1</sub>	809.344	62.5	–5.30 ± 0.02 –6.55 ± 0.05	3.48 ± 0.05 10.96 ± 0.12	13.33 ± 0.30 27.63 ± 0.71	3.59 ± 0.04 2.37 ± 0.04

Table A.1. continued.

Transition	Frequency [GHz]	$E_{\text{up}}/k$ [K]	$V_{\text{lsr}}$ [km s <sup>-1</sup> ]	$\Delta V$ [km s <sup>-1</sup> ]	$\int T_{\text{mb}} dV$ [K km s <sup>-1</sup> ]	$T_{\text{peak}}$ [K]
C <sup>+</sup>						
<sup>2</sup> P <sub>3/2</sub> – <sup>2</sup> P <sub>1/2</sub>	1900.537	91.2	–16.04 ± 0.05	4.95 ± 0.24	30.59 ± 4.62	5.81 ± 0.65
			–5.42 ± 0.05	3.11 ± 0.12	29.51 ± 2.56	8.92 ± 0.51
			–2.05 ± 0.03	3.21 ± 0.09	44.82 ± 2.94	13.10 ± 0.58
			–19.48 ± 2.60	12.79 ± 2.79	17.55 ± 7.91	1.29 ± 0.38
			–6.64 ± 0.71	9.05 ± 0.75	33.74 ± 6.24	3.51 ± 0.33
			1.59 ± 0.04	1.75 ± 0.13	4.86 ± 0.65	2.61 ± 0.20
HCO <sup>+</sup>						
4–3*	356.734	42.8	–5.87 ± 0.01	4.18 ± 0.02	61.7 ± 0.5	13.87 ± 0.04
			–7.61 ± 0.04	9.46 ± 0.08	39.8 ± 0.9	3.96 ± 0.02
6–5	535.061	89.9	–5.73 ± 0.03	3.48 ± 0.10	8.5 ± 0.2	2.30 ± 0.14
			–6.65 ± 0.21	6.12 ± 0.37	4.0 ± 0.4	0.62 ± 0.15
7–6	624.208	119.8	–5.55 ± 0.04	3.15 ± 0.14	6.0 ± 0.2	1.80 ± 0.20
			–6.42 ± 0.20	5.15 ± 0.27	4.5 ± 0.3	0.82 ± 0.20
8–7	713.342	154.1	–5.61 ± 0.01	3.10 ± 0.04	5.4 ± 0.1	1.65 ± 0.02
			–6.62 ± 0.14	8.29 ± 0.38	2.2 ± 0.4	0.25 ± 0.02
9–8	802.458	192.6	–5.47 ± 0.03	3.27 ± 0.06	4.8 ± 0.1	1.37 ± 0.02
10–9	891.557	235.3	–5.47 ± 0.04	3.11 ± 0.09	3.4 ± 0.1	1.02 ± 0.03
11–10	980.637	282.4	–5.21 ± 0.10	3.52 ± 0.25	2.1 ± 0.3	0.56 ± 0.04
H <sup>13</sup> CO <sup>+</sup>						
4–3*	346.998	41.6	–5.60 ± 0.10	2.90 ± 0.10	5.74 ± 0.36	1.86 ± 0.05
6–5	520.460	87.4	–5.49 ± 0.13	4.25 ± 0.31	0.66 ± 0.08	0.15 ± 0.01
7–6	607.175	116.6	–5.34 ± 0.15	1.96 ± 0.35	0.25 ± 0.08	0.12 ± 0.02
HCN						
4–3*	354.506	42.5	–5.51 ± 0.02	4.33 ± 0.04	36.20 ± 0.70	7.86 ± 0.28
			–6.52 ± 0.06	8.40 ± 0.20	22.40 ± 1.30	2.51 ± 0.30
6–5	531.716	89.3	–5.52 ± 0.02	3.91 ± 0.05	3.49 ± 0.08	0.84 ± 0.01
7–6	620.304	119.1	–5.35 ± 0.04	4.02 ± 0.09	2.88 ± 0.12	0.67 ± 0.02
8–7	708.877	153.1	–5.30 ± 0.06	4.16 ± 0.14	2.11 ± 0.13	0.48 ± 0.02
9–8	797.434	191.4	–5.14 ± 0.07	3.46 ± 0.17	1.20 ± 0.10	0.33 ± 0.02
10–9	885.971	233.9	–4.98 ± 0.09	5.47 ± 0.20	2.07 ± 0.13	0.36 ± 0.01
HCN $\nu = 1c, 4-3$	354.46043	1066.9	–5.22 ± 0.18	5.99 ± 0.45	2.29 ± 0.14	0.36 ± 0.10
HCN $\nu = 1d, 4-3$	356.25556	1067.1	–5.08 ± 0.25	5.27 ± 0.68	1.54 ± 0.15	0.28 ± 0.07
H <sup>13</sup> CN						
4–3*	345.340	41.4	–5.10 ± 0.10	5.40 ± 0.20	7.93 ± 0.44	1.38 ± 0.03
6–5	517.970	87.0	–5.44 ± 0.15	3.54 ± 0.36	0.46 ± 0.08	0.12 ± 0.02
7–6	604.268	116.0	–4.82 ± 0.14	3.38 ± 0.33	0.56 ± 0.09	0.16 ± 0.01
8–7	690.552	149.2	–4.92 ± 0.18	4.43 ± 0.44	0.64 ± 0.08	0.11 ± 0.01
HC <sup>15</sup> N						
4–3*	344.200	41.3	–5.50 ± 0.20	4.30 ± 0.40	2.52 ± 0.30	0.55 ± 0.04
6–5	516.262	86.7	–4.17 ± 0.23	3.66 ± 0.55	0.32 ± 0.08	0.08 ± 0.01
HNC						
4–3*	362.630	43.5	–5.51 ± 0.03	2.50 ± 0.10	8.20 ± 0.70	3.08 ± 0.13
			–6.01 ± 0.07	4.80 ± 0.20	10.00 ± 2.00	1.96 ± 0.12
6–5	543.897	91.4	–5.37 ± 0.06	3.36 ± 0.14	0.86 ± 0.06	0.24 ± 0.01
7–6	634.511	121.8	–5.32 ± 0.08	3.55 ± 0.19	0.79 ± 0.07	0.21 ± 0.01
CCH						
4 <sub>5</sub> –3 <sub>4</sub> *	349.338	41.9	–5.89 ± 0.04	3.80 ± 0.11	8.50 ± 0.19	2.12 ± 0.06
4 <sub>5</sub> –3 <sub>2</sub> *	349.401	41.9	–5.46 ± 0.32	4.01 ± 0.99	6.68 ± 1.21	1.56 ± 0.31
6 <sub>6</sub> –5 <sub>5</sub>	523.971	88.0	–6.41 ± 0.09	3.79 ± 0.21	0.85 ± 0.08	0.21 ± 0.01
6 <sub>5</sub> –5 <sub>4</sub>	524.033	88.0	–6.04 ± 0.11	4.13 ± 0.27	0.71 ± 0.08	0.16 ± 0.01
7 <sub>7</sub> –6 <sub>6</sub>	611.265	117.4	–6.26 ± 0.15	3.22 ± 0.35	0.47 ± 0.09	0.14 ± 0.01
			0.82 ± 0.15	2.58 ± 0.36	–0.33 ± 0.08	–0.12 ± 0.02
7 <sub>6</sub> –6 <sub>5</sub>	611.328	117.4	–6.64 ± 0.14	2.86 ± 0.33	0.39 ± 0.09	0.14 ± 0.02
8 <sub>8</sub> –7 <sub>7</sub>	698.542	150.9	–6.89 ± 0.19	3.89 ± 0.46	0.35 ± 0.08	0.09 ± 0.01
8 <sub>7</sub> –7 <sub>6</sub>	698.604	150.9	–6.64 ± 0.19	3.62 ± 0.45	0.38 ± 0.08	0.09 ± 0.01

Table A.1. continued.

Transition	Frequency [GHz]	$E_{\text{up}}/k$ [K]	$V_{\text{lsr}}$ [km s <sup>-1</sup> ]	$\Delta V$ [km s <sup>-1</sup> ]	$\int T_{\text{mb}} dV$ [K km s <sup>-1</sup> ]	$T_{\text{peak}}$ [K]
CH <sup>2</sup> Π <sub>3/2</sub>						
3/2 <sub>2+</sub> -1/2 <sub>1-</sub>	532.724	25.7	-5.10 ± 0.07	4.84 ± 0.19	2.00 ± 0.13	0.39 ± 0.01
			-0.32 ± 0.14	1.80 ± 0.33	-0.24 ± 0.07	-0.13 ± 0.02
3/2 <sub>1+</sub> -1/2 <sub>0-</sub>	532.793	25.7	-5.77 ± 0.14	4.83 ± 0.34	1.12 ± 0.13	0.22 ± 0.01
3/2 <sub>2-</sub> -1/2 <sub>1+</sub>	536.761	25.8	-5.40 ± 0.08	5.68 ± 0.20	2.04 ± 0.12	0.34 ± 0.01
			0.73 ± 0.15	2.57 ± 0.36	-0.31 ± 0.07	-0.11 ± 0.01
3/2 <sub>1-</sub> -1/2 <sub>1+</sub>	536.782	25.8	-4.93 ± 0.25	3.64 ± 0.60	1.98 ± 0.10	0.09 ± 0.01
3/2 <sub>1-</sub> -1/2 <sub>0+</sub>	536.796	25.8	-5.68 ± 0.15	4.92 ± 0.38	1.18 ± 0.12	0.19 ± 0.01
CH <sup>+</sup>						
1-0	835.138	40.1	4.17 ± 0.16	12.42 ± 0.43	-5.90 ± 0.33	-0.45 ± 0.01
			-7.46 ± 0.12	3.50 ± 0.29	1.24 ± 0.17	0.33 ± 0.02
			-16.90 ± 0.22	9.24 ± 0.59	-2.72 ± 0.28	-0.28 ± 0.01
OH <sup>2</sup> Π <sub>1/2</sub>						
$J = 3/2-1/2, F = 2--1+$	1834.747	269.8	-4.30 ± 0.18	3.42 ± 0.10	6.8 ± 0.3	1.27 ± 0.05
			-5.12 ± 0.06	10.39 ± 0.28	9.5 ± 0.6	0.56 ± 0.05
OH <sup>+</sup> , $F = 1-0$						
$J = 2-1, F = 3/2-1/2$	971.805	46.7	3.46 ± 0.28	11.87 ± 0.69	-7.66 ± 0.77	-0.61 ± 0.03
$J = 1-1, F = 3/2-1/2$	1033.004	49.6	3.88 ± 0.19	4.60 ± 0.47	-2.03 ± 0.36	-0.42 ± 0.04
$J = 1-1, F = 3/2-3/2$	1033.119	49.6	3.61 ± 0.25	10.92 ± 0.64	-5.78 ± 0.59	-0.51 ± 0.03
CN						
3 <sub>03</sub> -2 <sub>03</sub> *	339.517	32.6	-5.66 ± 0.30	3.19 ± 0.80	0.67 ± 0.17	0.20 ± 0.02
3 <sub>03</sub> -2 <sub>02</sub> *	340.008	32.6	-5.47 ± 0.18	2.14 ± 0.34	0.83 ± 0.35	0.36 ± 0.04
3 <sub>03</sub> -2 <sub>02</sub> *	340.020	32.6	-5.45 ± 0.37	3.14 ± 0.50	1.15 ± 0.31	0.34 ± 0.05
3 <sub>03</sub> -2 <sub>02</sub> *	340.032	32.6	-5.47 ± 0.19	2.93 ± 0.46	5.77 ± 1.01	1.85 ± 0.11
3 <sub>03</sub> -2 <sub>02</sub> *	340.035	32.6	-5.60 ± 0.20	3.17 ± 0.57	5.90 ± 1.04	1.75 ± 0.12
5 <sub>05</sub> -4 <sub>04</sub>	566.731	81.6	-5.61 ± 0.09	3.56 ± 0.22	0.88 ± 0.09	0.23 ± 0.01
5 <sub>06</sub> -4 <sub>05</sub>	566.947	81.7	-5.66 ± 0.07	3.52 ± 0.16	1.11 ± 0.09	0.30 ± 0.01
6 <sub>06</sub> -5 <sub>05</sub>	680.047	114.2	-5.38 ± 0.09	2.87 ± 0.21	0.47 ± 0.06	0.16 ± 0.01
6 <sub>07</sub> -5 <sub>06</sub>	680.264	114.3	-5.85 ± 0.11	3.80 ± 0.25	0.75 ± 0.07	0.17 ± 0.01
CS						
7-6*	342.883	65.8	-5.80 ± 0.02	3.10 ± 0.06	15.50 ± 0.50	4.70 ± 0.30
			-7.30 ± 0.30	6.40 ± 0.30	5.00 ± 0.80	0.73 ± 0.29
10-9	489.751	129.3	-5.68 ± 0.06	3.69 ± 0.14	1.30 ± 0.10	0.33 ± 0.01
11-10	538.689	155.2	-5.71 ± 0.07	3.53 ± 0.17	1.01 ± 0.20	0.27 ± 0.01
12-11	587.616	183.4	-5.04 ± 0.11	4.52 ± 0.25	0.89 ± 0.30	0.18 ± 0.01
13-12	636.532	213.9	-5.24 ± 0.12	4.04 ± 0.29	0.61 ± 0.30	0.14 ± 0.01
14-13	685.435	246.8	-5.05 ± 0.12	3.93 ± 0.28	0.73 ± 0.30	0.18 ± 0.01
15-14	734.324	282.0	-5.66 ± 0.16	4.17 ± 0.38	0.70 ± 0.40	0.16 ± 0.01
<sup>13</sup> CS						
8-7*	369.909	79.9	-5.20 ± 0.40	3.20 ± 0.80	0.85 ± 0.36	0.25 ± 0.05
C <sup>34</sup> S						
7-6*	337.396	50.2	-5.00 ± 0.30	3.40 ± 0.60	1.52 ± 0.46	0.42 ± 0.06
OCS						
28-27*	340.449	237.0	-4.91 ± 0.21	2.94 ± 0.40	0.77 ± 0.11	0.25 ± 0.03
29-28*	352.600	253.9	-4.85 ± 0.18	4.09 ± 0.44	1.12 ± 0.11	0.25 ± 0.03
30-29*	364.749	271.4	-5.37 ± 0.24	2.79 ± 0.80	0.80 ± 0.16	0.27 ± 0.04
o-H <sub>2</sub> S						
3 <sub>2,1</sub> -3 <sub>1,2</sub> *	369.102	154.5	-5.27 ± 0.32	3.30 ± 0.58	1.29 ± 0.22	0.37 ± 0.04
2 <sub>2,1</sub> -2 <sub>1,2</sub>	505.565	79.4	-5.71 ± 0.08	3.10 ± 0.20	0.70 ± 0.08	0.22 ± 0.01
2 <sub>1,2</sub> -1 <sub>0,1</sub>	736.034	55.1	-5.40 ± 0.03	3.39 ± 0.07	4.00 ± 0.14	1.11 ± 0.02
			0.22 ± 0.09	0.97 ± 0.22	-0.19 ± 0.08	-0.19 ± 0.04
3 <sub>03</sub> -2 <sub>1,2</sub>	993.102	350.1	-6.98 ± 0.16	4.34 ± 0.40	2.66 ± 0.42	0.58 ± 0.05
2 <sub>2,1</sub> -1 <sub>1,0</sub>	1072.837	79.4	-5.59 ± 0.19	4.05 ± 0.45	1.73 ± 0.30	0.38 ± 0.04
p-H <sub>2</sub> S						
3 <sub>3,1</sub> -3 <sub>2,2</sub>	568.051	166.0	-4.96 ± 0.19	2.08 ± 0.45	0.19 ± 0.07	0.08 ± 0.02
2 <sub>0,2</sub> -1 <sub>1,1</sub>	687.304	54.7	-5.77 ± 0.05	3.49 ± 0.13	1.43 ± 0.09	0.39 ± 0.01
H <sub>2</sub> <sup>34</sup> S						
2 <sub>1,2</sub> -1 <sub>0,1</sub>	734.269	55.0	-5.54 ± 0.18	1.94 ± 0.42	0.27 ± 0.12	0.15 ± 0.03

Table A.1. continued.

Transition	Frequency [GHz]	$E_{\text{up}}/k$ [K]	$V_{\text{lsr}}$ [km s <sup>-1</sup> ]	$\Delta V$ [km s <sup>-1</sup> ]	$\int T_{\text{mb}} dV$ [K km s <sup>-1</sup> ]	$T_{\text{peak}}$ [K]
o-H <sub>2</sub> CS						
10 <sub>1,10</sub> –9 <sub>1,9</sub> *	338.083	102.4	$-5.60 \pm 0.27$	$3.09 \pm 1.07$	$0.96 \pm 0.21$	$0.30 \pm 0.03$
10 <sub>1,9</sub> –9 <sub>1,8</sub> *	348.534	105.2	$-5.25 \pm 0.29$	$4.61 \pm 0.86$	$1.31 \pm 0.18$	$0.27 \pm 0.03$
11 <sub>1,11</sub> –10 <sub>1,10</sub> *	371.847	120.3	$-5.62 \pm 0.33$	$4.36 \pm 0.71$	$0.98 \pm 0.15$	$0.21 \pm 0.03$
HCl						
1 <sub>3/2</sub> –0 <sub>3/2</sub>	625.902	30.0	$-5.85 \pm 0.04$	$3.63 \pm 0.08$	$2.46 \pm 0.10$	$0.64 \pm 0.01$
1 <sub>5/2</sub> –0 <sub>3/2</sub>	625.919	30.0	$-5.68 \pm 0.03$	$3.82 \pm 0.08$	$2.45 \pm 0.10$	$0.73 \pm 0.01$
1 <sub>1/2</sub> –0 <sub>3/2</sub>	625.932	30.0	$-5.89 \pm 0.06$	$3.89 \pm 0.14$	$2.44 \pm 0.10$	$0.42 \pm 0.01$
2 <sub>5/2</sub> –1 <sub>5/2</sub>	1251.434	90.1	$-5.21 \pm 0.12$	$2.22 \pm 0.37$	$0.52 \pm 0.28$	$0.30 \pm 0.08$
2 <sub>7/2</sub> –1 <sub>5/2</sub>	1251.452	90.1	$-5.27 \pm 0.03$	$4.51 \pm 0.08$	$2.26 \pm 0.20$	$0.47 \pm 0.04$
H <sup>37</sup> Cl						
1 <sub>3/2</sub> –0 <sub>3/2</sub>	624.964	30.0	$-5.74 \pm 0.09$	$4.07 \pm 0.23$	$1.15 \pm 0.11$	$0.27 \pm 0.01$
1 <sub>5/2</sub> –0 <sub>3/2</sub>	624.978	30.0	$-5.86 \pm 0.07$	$3.19 \pm 0.18$	$1.18 \pm 0.12$	$0.42 \pm 0.02$
1 <sub>1/2</sub> –0 <sub>3/2</sub>	624.988	30.0	$-5.74 \pm 0.18$	$3.89 \pm 0.45$	$0.17 \pm 0.13$	$0.17 \pm 0.01$
o-NH <sub>3</sub>						
1 <sub>0</sub> –0 <sub>0</sub>	572.498	27.5	$-5.41 \pm 0.03$	$4.13 \pm 0.07$	$3.41 \pm 0.10$	$0.78 \pm 0.01$
			$0.00 \pm 0.15$	$1.30 \pm 0.36$	$-0.12 \pm 0.06$	$-0.09 \pm 0.02$
3 <sub>0</sub> –2 <sub>0</sub>	1763.524	170.4	$-5.30 \pm 0.13$	$5.41 \pm 0.31$	$2.54 \pm 0.25$	$0.44 \pm 0.02$
p-NH <sub>3</sub>						
2 <sub>1</sub> –1 <sub>1</sub>	1168.452	79.3	$-5.68 \pm 0.11$	$2.68 \pm 0.28$	$2.29 \pm 0.41$	$0.80 \pm 0.07$
3 <sub>1</sub> –2 <sub>1</sub>	1763.601	165.1	$-4.37 \pm 0.09$	$4.18 \pm 0.22$	$1.01 \pm 0.32$	$0.23 \pm 0.07$
3 <sub>2</sub> –2 <sub>2</sub>	1763.823	149.1	$-4.55 \pm 0.15$	$3.92 \pm 0.36$	$1.67 \pm 0.25$	$0.40 \pm 0.03$
N <sub>2</sub> H <sup>+</sup>						
4–3*	372.673	44.7	$-6.00 \pm 0.10$	$2.90 \pm 0.10$	$6.61 \pm 0.39$	$2.14 \pm 0.06$
6–5	558.967	93.9	$-5.68 \pm 0.09$	$2.89 \pm 0.21$	$0.76 \pm 0.09$	$0.25 \pm 0.02$
7–6	652.096	125.2	$-6.01 \pm 0.09$	$2.47 \pm 0.19$	$0.43 \pm 0.06$	$0.16 \pm 0.01$
NO						
<sup>2</sup> Π <sub>+1/2</sub> , $J = 7/2-5/2$ , $F = 7/2-5/2^*$	350.691	36.1	$-4.40 \pm 0.20$	$4.90 \pm 0.30$	$4.32 \pm 0.13$	$0.80 \pm 0.10$
<sup>2</sup> Π <sub>-1/2</sub> , $J = 7/2-5/2$ , $F = 7/2-5/2^*$	351.052	36.1	$-4.62 \pm 0.70$	$9.98 \pm 1.50$	$2.01 \pm 0.14$	$0.17 \pm 0.10$
<sup>2</sup> Π <sub>+1/2</sub> , $J=13/2-11/2$ , $F=11/2-9/2$	651.434	115.4	$-5.39 \pm 0.17$	$4.46 \pm 0.40$	$0.48 \pm 0.08$	$0.10 \pm 0.01$
<sup>2</sup> Π <sub>-1/2</sub> , $J=13/2-11/2$ , $F=11/2-9/2$	651.773	115.5	$-4.92 \pm 0.09$	$3.78 \pm 0.22$	$0.58 \pm 0.06$	$0.15 \pm 0.01$
E-CH <sub>3</sub> OH						
7 <sub>0,0</sub> –6 <sub>-1,0</sub>	495.173	70.2	$-5.62 \pm 0.20$	$3.16 \pm 0.47$	$0.35 \pm 0.10$	$0.11 \pm 0.02$
7 <sub>1,0</sub> –6 <sub>0,0</sub>	504.294	78.2	$-6.25 \pm 0.19$	$4.10 \pm 0.46$	$0.49 \pm 0.10$	$0.11 \pm 0.01$
2 <sub>-2,0</sub> –1 <sub>-1,0</sub>	520.179	25.0	$-5.33 \pm 0.13$	$3.57 \pm 0.30$	$0.57 \pm 0.09$	$0.15 \pm 0.01$
13 <sub>-4,0</sub> –13 <sub>-3,0</sub>	524.269	291.2	$-5.86 \pm 0.18$	$3.61 \pm 0.44$	$0.40 \pm 0.08$	$0.10 \pm 0.01$
11 <sub>2,0</sub> –10 <sub>1,0</sub>	558.345	167.6	$-5.57 \pm 0.24$	$3.10 \pm 0.57$	$0.34 \pm 0.11$	$0.10 \pm 0.02$
3 <sub>-2,0</sub> –2 <sub>-1,0</sub>	568.566	31.9	$-5.87 \pm 0.17$	$4.08 \pm 0.41$	$0.58 \pm 0.10$	$0.14 \pm 0.01$
12 <sub>-2,0</sub> –11 <sub>-2,0</sub>	581.092	199.2	$-6.21 \pm 0.20$	$2.55 \pm 0.46$	$0.23 \pm 0.07$	$0.09 \pm 0.01$
13 <sub>1,0</sub> –12 <sub>2,0</sub>	601.849	224.4	$-6.24 \pm 0.25$	$3.44 \pm 0.60$	$0.30 \pm 0.09$	$0.08 \pm 0.01$
9 <sub>1,0</sub> –8 <sub>0,0</sub>	602.233	117.6	$-5.86 \pm 0.21$	$2.86 \pm 0.49$	$0.33 \pm 0.10$	$0.11 \pm 0.02$
4 <sub>-2,0</sub> –3 <sub>-1,0</sub>	616.980	41.2	$-5.63 \pm 0.11$	$3.39 \pm 0.27$	$0.62 \pm 0.09$	$0.17 \pm 0.01$
13 <sub>0,0</sub> –12 <sub>0,0</sub>	625.749	215.9	$-6.31 \pm 0.31$	$4.52 \pm 0.74$	$0.38 \pm 0.11$	$0.08 \pm 0.01$
13 <sub>-2,0</sub> –12 <sub>-2,0</sub>	629.652	229.4	$-6.56 \pm 0.23$	$3.21 \pm 0.55$	$0.30 \pm 0.09$	$0.09 \pm 0.01$
10 <sub>0,0</sub> –9 <sub>-1,0</sub>	638.280	132.7	$-5.53 \pm 0.18$	$2.98 \pm 0.42$	$0.31 \pm 0.08$	$0.10 \pm 0.01$
14 <sub>1,0</sub> –13 <sub>2,0</sub>	649.540	256.9	$-5.84 \pm 0.17$	$3.71 \pm 0.42$	$0.27 \pm 0.05$	$0.07 \pm 0.01$
10 <sub>1,0</sub> –9 <sub>0,0</sub>	651.617	140.8	$-5.79 \pm 0.12$	$3.79 \pm 0.30$	$0.40 \pm 0.06$	$0.10 \pm 0.01$
5 <sub>-2,0</sub> –4 <sub>-1,0</sub>	665.442	52.8	$-5.53 \pm 0.07$	$2.87 \pm 0.16$	$0.54 \pm 0.06$	$0.20 \pm 0.01$
17 <sub>-1,0</sub> –16 <sub>0,0</sub>	672.903	352.0	$-5.76 \pm 0.14$	$3.09 \pm 0.33$	$0.35 \pm 0.06$	$0.11 \pm 0.01$
3 <sub>3,0</sub> –2 <sub>2,0</sub>	675.773	53.7	$-5.81 \pm 0.20$	$4.17 \pm 0.49$	$0.44 \pm 0.08$	$0.09 \pm 0.01$
11 <sub>0,0</sub> –10 <sub>-1,0</sub>	685.505	158.2	$-5.56 \pm 0.23$	$4.30 \pm 0.56$	$0.50 \pm 0.08$	$0.09 \pm 0.01$
14 <sub>2,0</sub> –13 <sub>1,0</sub>	705.182	258.2	$-5.11 \pm 0.16$	$2.61 \pm 0.38$	$0.23 \pm 0.06$	$0.08 \pm 0.01$
6 <sub>-2,0</sub> –5 <sub>-1,0</sub>	713.983	66.8	$-5.93 \pm 0.10$	$3.47 \pm 0.22$	$0.63 \pm 0.07$	$0.17 \pm 0.01$
15 <sub>0,0</sub> –14 <sub>0,0</sub>	721.011	282.9	$-6.11 \pm 0.14$	$2.64 \pm 0.33$	$0.44 \pm 0.10$	$0.16 \pm 0.02$
13 <sub>1,0</sub> –12 <sub>0,0</sub>	802.241	224.4	$-5.78 \pm 0.13$	$2.14 \pm 0.31$	$0.43 \pm 0.11$	$0.19 \pm 0.02$
6 <sub>-4,0</sub> –5 <sub>-3,0</sub>	815.071	128.8	$-5.41 \pm 0.12$	$2.63 \pm 0.28$	$0.43 \pm 0.08$	$0.15 \pm 0.01$
6 <sub>3,0</sub> –5 <sub>2,0</sub>	820.762	88.6	$-5.83 \pm 0.22$	$3.45 \pm 0.52$	$0.46 \pm 0.12$	$0.13 \pm 0.02$



Table A.1. continued.

Transition	Frequency [GHz]	$E_{\text{up}}/k$ [K]	$V_{\text{lsr}}$ [km s <sup>-1</sup> ]	$\Delta V$ [km s <sup>-1</sup> ]	$\int T_{\text{mb}} dV$ [K km s <sup>-1</sup> ]	$T_{\text{peak}}$ [K]
A-CH <sub>3</sub> OH						
4 <sub>1,+0</sub> -3 <sub>0,+0</sub>	492.279	37.6	-5.75 ± 0.11	3.44 ± 0.26	0.80 ± 0.11	0.22 ± 0.01
5 <sub>3,+0</sub> -4 <sub>2,+0</sub>	493.699	84.6	-5.41 ± 0.17	2.67 ± 0.41	0.30 ± 0.08	0.11 ± 0.01
5 <sub>1,+0</sub> -4 <sub>0,+0</sub>	538.571	49.1	-5.76 ± 0.07	3.03 ± 0.16	0.72 ± 0.07	0.22 ± 0.01
6 <sub>3,+0</sub> -5 <sub>2,+0</sub>	542.001	98.6	-5.60 ± 0.14	1.88 ± 0.33	0.31 ± 0.10	0.16 ± 0.02
6 <sub>3,-0</sub> -5 <sub>2,-0</sub>	542.082	98.6	-6.69 ± 0.19	3.43 ± 0.47	0.60 ± 0.14	0.16 ± 0.02
2 <sub>2,-0</sub> -1 <sub>1,-0</sub>	579.085	44.7	-5.44 ± 0.11	3.01 ± 0.27	0.54 ± 0.09	0.17 ± 0.01
2 <sub>2,+0</sub> -1 <sub>1,+0</sub>	579.921	44.7	-6.39 ± 0.21	4.30 ± 0.50	0.60 ± 0.13	0.13 ± 0.01
12 <sub>4,+0</sub> -11 <sub>4,+0</sub>	580.196	261.4	-5.87 ± 0.20	2.57 ± 0.49	0.20 ± 0.09	0.08 ± 0.02
12 <sub>3,-0</sub> -11 <sub>3,-0</sub>	580.213	230.8	-5.51 ± 0.26	3.53 ± 0.62	0.31 ± 0.10	0.09 ± 0.01
6 <sub>1,+0</sub> -5 <sub>0,+0</sub>	584.450	62.9	-5.54 ± 0.12	3.07 ± 0.27	0.66 ± 0.10	0.20 ± 0.02
13 <sub>1,+0</sub> -12 <sub>1,+0</sub>	622.659	223.8	-6.25 ± 0.18	3.58 ± 0.43	0.40 ± 0.09	0.10 ± 0.01
3 <sub>2,-0</sub> -2 <sub>1,-0</sub>	626.626	51.6	-6.25 ± 0.13	3.67 ± 0.31	0.71 ± 0.10	0.18 ± 0.01
7 <sub>1,+0</sub> -6 <sub>0,+0</sub>	629.921	79.0	-5.32 ± 0.11	3.16 ± 0.11	0.65 ± 0.10	0.19 ± 0.01
13 <sub>1,-0</sub> -12 <sub>1,-0</sub>	633.423	227.5	-6.74 ± 0.24	4.14 ± 0.59	0.50 ± 0.12	0.11 ± 0.01
8 <sub>3,-0</sub> -7 <sub>2,-0</sub>	638.818	133.4	-5.41 ± 0.13	3.13 ± 0.31	0.51 ± 0.09	0.15 ± 0.01
4 <sub>2,-0</sub> -3 <sub>1,-0</sub>	673.746	60.9	-5.70 ± 0.09	3.66 ± 0.22	0.69 ± 0.07	0.18 ± 0.01
8 <sub>1,+0</sub> -7 <sub>0,+0</sub>	674.991	97.4	-5.42 ± 0.09	3.18 ± 0.22	0.56 ± 0.07	0.17 ± 0.01
14 <sub>4,+0</sub> -13 <sub>4,+0</sub>	676.824	324.0	-5.48 ± 0.15	3.89 ± 0.36	0.47 ± 0.07	0.11 ± 0.01
4 <sub>2,+0</sub> -3 <sub>1,+0</sub>	678.785	60.9	-5.99 ± 0.09	4.13 ± 0.20	0.87 ± 0.07	0.20 ± 0.01
9 <sub>3,+0</sub> -8 <sub>2,+0</sub>	686.732	154.3	-5.36 ± 0.15	4.20 ± 0.37	0.54 ± 0.08	0.12 ± 0.01
9 <sub>3,-0</sub> -8 <sub>2,-0</sub>	687.225	154.3	-5.52 ± 0.23	2.70 ± 0.55	0.32 ± 0.11	0.11 ± 0.02
5 <sub>2,+0</sub> -4 <sub>1,+0</sub>	728.863	72.5	-5.46 ± 0.18	4.19 ± 0.43	0.95 ± 0.17	0.21 ± 0.02
16 <sub>0,+0</sub> -15 <sub>0,+0</sub>	771.576	315.2	-4.57 ± 0.19	3.21 ± 0.44	1.70 ± 0.31	0.41 ± 0.04
4 <sub>4,-0</sub> -3 <sub>3,-0</sub>	829.891	103.6	-5.26 ± 0.14	3.40 ± 0.36	0.67 ± 0.11	0.18 ± 0.02
7 <sub>2,+0</sub> -6 <sub>1,+0</sub>	830.349	102.7	-5.60 ± 0.12	2.67 ± 0.29	0.57 ± 0.11	0.20 ± 0.02
12 <sub>3,-0</sub> -11 <sub>2,-0</sub>	832.754	230.8	-5.45 ± 0.16	2.73 ± 0.38	0.49 ± 0.12	0.17 ± 0.02
13 <sub>1,+0</sub> -12 <sub>0,+0</sub>	894.614	223.8	-6.08 ± 0.18	3.59 ± 0.43	0.71 ± 0.15	0.19 ± 0.02
9 <sub>2,-0</sub> -8 <sub>1,-0</sub>	902.935	142.2	-5.45 ± 0.22	3.12 ± 0.52	0.62 ± 0.18	0.19 ± 0.03
9 <sub>4,+0</sub> -8 <sub>3,+0</sub>	1071.514	184.8	-4.69 ± 0.10	1.95 ± 0.23	0.97 ± 0.19	0.47 ± 0.05
o-H <sub>2</sub> CO						
5 <sub>15</sub> -4 <sub>14</sub> *	351.769	31.7	-5.57 ± 0.02	3.51 ± 0.04	10.85 ± 0.10	2.91 ± 0.02
5 <sub>33</sub> -4 <sub>32</sub> *	364.275	158.4	-5.24 ± 0.16	4.78 ± 0.35	2.98 ± 0.21	0.58 ± 0.08
5 <sub>32</sub> -4 <sub>31</sub> *	364.289	158.4	-5.65 ± 0.26	4.21 ± 0.63	2.96 ± 0.38	0.66 ± 0.17
7 <sub>17</sub> -6 <sub>16</sub>	491.968	106.3	-5.75 ± 0.09	3.59 ± 0.21	0.98 ± 0.10	0.26 ± 0.01
7 <sub>35</sub> -6 <sub>34</sub>	510.156	203.9	-5.21 ± 0.56	3.19 ± 0.97	0.33 ± 0.15	0.10 ± 0.02
7 <sub>34</sub> -6 <sub>33</sub>	510.238	203.9	-5.43 ± 0.25	3.64 ± 0.60	0.31 ± 0.09	0.08 ± 0.01
7 <sub>16</sub> -6 <sub>15</sub>	525.666	112.8	-5.56 ± 0.09	3.56 ± 0.25	0.77 ± 0.09	0.20 ± 0.01
8 <sub>18</sub> -7 <sub>17</sub>	561.899	133.3	-4.69 ± 0.12	3.53 ± 0.28	0.80 ± 0.11	0.21 ± 0.02
8 <sub>17</sub> -7 <sub>16</sub>	600.331	141.6	-5.20 ± 0.15	3.53 ± 0.36	0.61 ± 0.11	0.16 ± 0.02
9 <sub>19</sub> -8 <sub>18</sub>	631.703	163.6	-5.09 ± 0.12	2.90 ± 0.28	0.53 ± 0.09	0.17 ± 0.01
9 <sub>36</sub> -8 <sub>35</sub>	656.465	263.4	-4.84 ± 0.22	4.43 ± 0.54	0.36 ± 0.05	0.08 ± 0.01
9 <sub>18</sub> -8 <sub>17</sub>	674.810	174.0	-5.18 ± 0.22	4.49 ± 0.52	0.49 ± 0.05	0.10 ± 0.01
p-H <sub>2</sub> CO						
5 <sub>05</sub> -4 <sub>04</sub> *	362.736	52.3	-5.58 ± 0.04	3.47 ± 0.10	7.27 ± 0.16	1.96 ± 0.04
5 <sub>24</sub> -4 <sub>23</sub> *	363.946	99.5	-5.57 ± 0.21	3.85 ± 0.47	2.48 ± 0.27	0.61 ± 0.13
5 <sub>23</sub> -4 <sub>22</sub> *	365.363	99.7	-5.69 ± 0.10	3.53 ± 0.28	1.98 ± 0.12	0.53 ± 0.09
7 <sub>07</sub> -6 <sub>06</sub>	505.834	97.4	-5.53 ± 0.09	1.92 ± 0.20	0.36 ± 0.07	0.18 ± 0.02
7 <sub>25</sub> -6 <sub>24</sub>	513.076	145.4	-5.30 ± 0.17	2.29 ± 0.39	0.23 ± 0.07	0.09 ± 0.01

Table A.1. continued.

Transition	Frequency [GHz]	$E_{\text{up}}/k$ [K]	$V_{\text{lsr}}$ [km s <sup>-1</sup> ]	$\Delta V$ [km s <sup>-1</sup> ]	$\int T_{\text{mb}} dV$ [K km s <sup>-1</sup> ]	$T_{\text{peak}}$ [K]
SO						
3 <sub>3</sub> –3 <sub>2</sub> *	339.342	25.5	-5.39 ± 0.12	3.38 ± 0.32	1.58 ± 0.12	0.44 ± 0.05
7 <sub>8</sub> –6 <sub>7</sub> *	340.714	81.2	-5.78 ± 0.07	5.35 ± 0.17	7.99 ± 0.20	1.41 ± 0.10
8 <sub>8</sub> –7 <sub>7</sub> *	344.311	87.5	-5.76 ± 0.06	5.12 ± 0.13	8.57 ± 0.18	1.57 ± 0.10
9 <sub>8</sub> –8 <sub>7</sub> *	346.529	78.8	-5.56 ± 0.05	5.91 ± 0.14	10.42 ± 0.18	1.66 ± 0.10
12 <sub>11</sub> –11 <sub>10</sub>	514.853	167.6	-5.43 ± 0.18	4.51 ± 0.44	0.61 ± 0.10	0.13 ± 0.01
12 <sub>12</sub> –11 <sub>11</sub>	516.335	174.2	-4.97 ± 0.23	5.65 ± 0.55	0.72 ± 0.12	0.12 ± 0.01
12 <sub>13</sub> –11 <sub>12</sub>	517.354	165.8	-5.08 ± 0.18	5.15 ± 0.43	0.73 ± 0.11	0.13 ± 0.01
13 <sub>12</sub> –12 <sub>11</sub>	558.087	194.4	-5.83 ± 0.18	5.50 ± 0.44	0.95 ± 0.13	0.16 ± 0.01
13 <sub>13</sub> –12 <sub>12</sub>	559.319	201.1	-4.90 ± 0.20	5.73 ± 0.48	0.90 ± 0.13	0.15 ± 0.01
13 <sub>14</sub> –12 <sub>13</sub>	560.178	192.7	-6.30 ± 0.15	4.47 ± 0.36	0.75 ± 0.11	0.16 ± 0.01
14 <sub>13</sub> –13 <sub>12</sub>	601.258	223.2	-5.90 ± 0.18	5.08 ± 0.44	0.79 ± 0.12	0.15 ± 0.01
14 <sub>14</sub> –13 <sub>13</sub>	602.292	230.0	-4.87 ± 0.27	4.56 ± 0.65	0.53 ± 0.13	0.11 ± 0.01
14 <sub>15</sub> –13 <sub>14</sub>	603.021	221.6	-5.60 ± 0.17	5.04 ± 0.40	0.81 ± 0.11	0.15 ± 0.01
15 <sub>14</sub> –14 <sub>13</sub>	644.378	254.2	-5.47 ± 0.13	4.61 ± 0.32	0.80 ± 0.10	0.16 ± 0.01
15 <sub>15</sub> –14 <sub>14</sub>	645.254	260.9	-5.18 ± 0.15	5.14 ± 0.37	0.80 ± 0.10	0.14 ± 0.01
15 <sub>16</sub> –14 <sub>15</sub>	645.875	252.6	-5.51 ± 0.12	4.86 ± 0.29	0.86 ± 0.09	0.16 ± 0.01
16 <sub>15</sub> –15 <sub>14</sub>	687.456	287.2	-5.45 ± 0.12	4.91 ± 0.29	0.82 ± 0.08	0.15 ± 0.01
16 <sub>16</sub> –15 <sub>15</sub>	688.204	294.0	-5.24 ± 0.10	3.39 ± 0.25	0.62 ± 0.09	0.17 ± 0.01
16 <sub>17</sub> –15 <sub>16</sub>	688.735	285.7	-5.72 ± 0.16	5.36 ± 0.38	0.96 ± 0.08	0.14 ± 0.01
19 <sub>18</sub> –18 <sub>17</sub>	816.493	398.5	-6.06 ± 0.19	3.88 ± 0.46	0.68 ± 0.09	0.16 ± 0.02
19 <sub>19</sub> –18 <sub>18</sub>	816.971	405.4	-6.01 ± 0.20	4.64 ± 0.48	0.76 ± 0.13	0.15 ± 0.01
19 <sub>20</sub> –18 <sub>19</sub>	817.306	397.2	-5.32 ± 0.19	4.41 ± 0.47	0.76 ± 0.14	0.16 ± 0.01
<sup>34</sup> SO						
7 <sub>8</sub> –6 <sub>7</sub> *	333.902	79.9	-5.12 ± 0.18	4.15 ± 0.43	1.74 ± 0.16	0.39 ± 0.05
8 <sub>8</sub> –7 <sub>7</sub> *	337.582	86.1	-5.41 ± 0.27	4.05 ± 0.62	1.12 ± 0.15	0.26 ± 0.04
9 <sub>8</sub> –8 <sub>7</sub> *	339.858	77.3	-5.18 ± 0.13	4.75 ± 0.30	2.06 ± 0.12	0.41 ± 0.05
SO <sub>2</sub>						
21 <sub>2,20</sub> –21 <sub>1,21</sub> *	332.091	219.5	-4.74 ± 0.18	5.00 ± 0.43	1.81 ± 0.14	0.34 ± 0.04
4 <sub>3,1</sub> –3 <sub>2,2</sub> *	332.505	31.3	-5.41 ± 0.14	5.26 ± 0.36	2.80 ± 0.16	0.50 ± 0.07
8 <sub>2,6</sub> –7 <sub>1,7</sub> *	334.673	43.2	-5.25 ± 0.17	5.43 ± 0.37	3.52 ± 0.22	0.61 ± 0.08
23 <sub>3,21</sub> –23 <sub>2,22</sub> *	336.089	276.0	-4.55 ± 0.21	4.44 ± 0.46	1.87 ± 0.18	0.40 ± 0.06
18 <sub>4,14</sub> –18 <sub>3,15</sub> *	338.306	196.8	-5.03 ± 0.19	5.29 ± 0.41	2.31 ± 0.16	0.41 ± 0.06
20 <sub>1,19</sub> –19 <sub>2,18</sub> *	338.612	198.9	-6.55 ± 0.34	6.04 ± 0.50	3.95 ± 0.46	0.61 ± 0.09
19 <sub>1,19</sub> –18 <sub>0,18</sub> *	346.652	168.1	-5.11 ± 0.14	5.99 ± 0.33	3.45 ± 0.16	0.54 ± 0.07
24 <sub>2,22</sub> –23 <sub>3,21</sub> *	348.388	292.7	-4.98 ± 0.18	5.06 ± 0.40	2.47 ± 0.18	0.46 ± 0.06
10 <sub>6,4</sub> –11 <sub>5,7</sub> *	350.863	138.9	-5.41 ± 0.20	4.15 ± 0.40	0.82 ± 0.10	0.19 ± 0.03
5 <sub>3,3</sub> –4 <sub>2,2</sub> *	351.257	35.9	-5.16 ± 0.10	5.16 ± 0.21	3.30 ± 0.12	0.60 ± 0.07
14 <sub>4,10</sub> –14 <sub>3,11</sub> *	351.874	135.9	-5.10 ± 0.10	4.90 ± 0.22	2.31 ± 0.10	0.44 ± 0.05
12 <sub>4,8</sub> –12 <sub>3,9</sub> *	355.046	111.0	-5.02 ± 0.10	5.29 ± 0.21	3.84 ± 0.13	0.68 ± 0.08
13 <sub>4,10</sub> –13 <sub>3,11</sub> *	357.165	122.9	-5.12 ± 0.16	5.27 ± 0.34	2.70 ± 0.16	0.48 ± 0.06
15 <sub>4,12</sub> –15 <sub>3,13</sub> *	357.241	149.7	-5.25 ± 0.11	5.10 ± 0.26	2.54 ± 0.12	0.47 ± 0.06
11 <sub>4,8</sub> –11 <sub>3,9</sub> *	357.388	100.0	-5.11 ± 0.11	5.12 ± 0.22	2.99 ± 0.12	0.55 ± 0.06
8 <sub>4,4</sub> –8 <sub>3,5</sub> *	357.581	72.4	-5.01 ± 0.13	4.64 ± 0.26	2.58 ± 0.14	0.52 ± 0.06
9 <sub>4,6</sub> –9 <sub>3,7</sub> *	357.672	80.6	-5.14 ± 0.13	5.23 ± 0.32	2.85 ± 0.15	0.51 ± 0.06
7 <sub>4,4</sub> –7 <sub>3,5</sub> *	357.892	65.0	-4.85 ± 0.10	4.71 ± 0.18	3.02 ± 0.10	0.60 ± 0.06
6 <sub>4,2</sub> –6 <sub>3,3</sub> *	357.926	58.6	-5.24 ± 0.26	5.65 ± 0.63	2.98 ± 0.28	0.49 ± 0.06
4 <sub>4,0</sub> –4 <sub>3,1</sub> *	358.038	48.5	-5.32 ± 0.12	4.36 ± 0.25	2.07 ± 0.11	0.45 ± 0.05
20 <sub>0,20</sub> –19 <sub>1,19</sub> *	358.216	185.3	-4.97 ± 0.10	5.26 ± 0.22	3.78 ± 0.13	0.68 ± 0.08
21 <sub>4,18</sub> –21 <sub>3,19</sub> *	363.159	252.1	-5.39 ± 0.20	5.47 ± 0.48	2.62 ± 0.20	0.45 ± 0.06
15 <sub>2,14</sub> –14 <sub>1,13</sub> *	366.215	119.3	-5.18 ± 0.10	5.09 ± 0.23	3.17 ± 0.13	0.58 ± 0.06
6 <sub>3,3</sub> –5 <sub>2,4</sub> *	371.172	41.1	-5.19 ± 0.13	4.86 ± 0.29	2.75 ± 0.14	0.53 ± 0.06
28 <sub>0,28</sub> –27 <sub>1,27</sub>	501.108	354.3	-5.41 ± 0.17	3.12 ± 0.42	0.38 ± 0.09	0.11 ± 0.01
15 <sub>3,13</sub> –14 <sub>2,12</sub>	508.710	132.5	-4.63 ± 0.19	3.40 ± 0.46	0.41 ± 0.09	0.11 ± 0.01
18 <sub>6,12</sub> –18 <sub>5,13</sub>	559.882	245.5	-5.88 ± 0.24	3.06 ± 0.57	0.32 ± 0.11	0.10 ± 0.02
13 <sub>6,8</sub> –13 <sub>5,9</sub>	561.266	171.9	-5.27 ± 0.25	4.81 ± 0.61	0.61 ± 0.12	0.11 ± 0.01
12 <sub>6,6</sub> –12 <sub>5,7</sub>	561.393	160.0	-5.13 ± 0.21	2.96 ± 0.49	0.32 ± 0.09	0.10 ± 0.02
8 <sub>5,3</sub> –7 <sub>4,4</sub>	613.076	94.4	-4.86 ± 0.19	4.74 ± 0.45	0.53 ± 0.09	0.11 ± 0.01
14 <sub>4,10</sub> –13 <sub>3,11</sub>	626.087	135.9	-4.54 ± 0.10	2.23 ± 0.24	0.41 ± 0.08	0.17 ± 0.02
9 <sub>5,5</sub> –8 <sub>4,4</sub>	632.193	102.7	-5.09 ± 0.20	4.61 ± 0.48	0.56 ± 0.10	0.11 ± 0.01

Table A.1. continued.

Transition	Frequency [GHz]	$E_{\text{up}}/k$ [K]	$V_{\text{lsr}}$ [km s <sup>-1</sup> ]	$\Delta V$ [km s <sup>-1</sup> ]	$\int T_{\text{mb}} dV$ [K km s <sup>-1</sup> ]	$T_{\text{peak}}$ [K]
15 <sub>4,12</sub> –14 <sub>3,11</sub>	639.651	149.7	$-5.16 \pm 0.20$	$3.76 \pm 0.48$	$0.37 \pm 0.08$	$0.09 \pm 0.01$
10 <sub>5,5</sub> –9 <sub>4,6</sub>	651.300	111.9	$-5.06 \pm 0.15$	$5.06 \pm 0.35$	$0.62 \pm 0.07$	$0.11 \pm 0.01$
18 <sub>3,15</sub> –17 <sub>2,16</sub>	653.110	180.6	$-4.83 \pm 0.18$	$5.01 \pm 0.44$	$0.55 \pm 0.08$	$0.10 \pm 0.01$
22 <sub>7,15</sub> –22 <sub>6,16</sub>	660.918	352.8	$-4.81 \pm 0.21$	$3.69 \pm 0.49$	$0.39 \pm 0.09$	$0.10 \pm 0.01$
19 <sub>7,13</sub> –19 <sub>6,14</sub>	661.962	294.8	$-5.00 \pm 0.18$	$3.60 \pm 0.44$	$0.35 \pm 0.08$	$0.09 \pm 0.01$
17 <sub>7,11</sub> –17 <sub>6,12</sub>	662.404	260.8	$-4.64 \pm 0.25$	$4.50 \pm 0.60$	$0.34 \pm 0.08$	$0.07 \pm 0.01$
16 <sub>7,9</sub> –16 <sub>6,10</sub>	662.567	245.1	$-4.99 \pm 0.15$	$3.91 \pm 0.36$	$0.42 \pm 0.07$	$0.10 \pm 0.01$
14 <sub>7,7</sub> –14 <sub>6,8</sub>	662.799	216.6	$-5.34 \pm 0.19$	$4.12 \pm 0.46$	$0.43 \pm 0.08$	$0.10 \pm 0.01$
13 <sub>7,7</sub> –14 <sub>6,8</sub>	662.877	203.8	$-5.19 \pm 0.23$	$4.73 \pm 0.57$	$0.36 \pm 0.08$	$0.07 \pm 0.01$
12 <sub>7,5</sub> –12 <sub>6,6</sub>	662.934	191.8	$-4.93 \pm 0.22$	$3.92 \pm 0.53$	$0.40 \pm 0.08$	$0.08 \pm 0.01$
16 <sub>4,12</sub> –15 <sub>3,13</sub>	665.247	164.5	$-4.75 \pm 0.21$	$5.53 \pm 0.49$	$0.59 \pm 0.08$	$0.10 \pm 0.01$
11 <sub>5,7</sub> –10 <sub>4,6</sub>	670.366	122.0	$-5.16 \pm 0.16$	$4.07 \pm 0.37$	$0.51 \pm 0.08$	$0.12 \pm 0.01$
7 <sub>6,2</sub> –6 <sub>5,1</sub>	695.633	114.0	$-4.89 \pm 0.13$	$3.73 \pm 0.30$	$0.51 \pm 0.07$	$0.13 \pm 0.01$
19 <sub>4,16</sub> –18 <sub>3,15</sub>	702.104	214.3	$-4.73 \pm 0.17$	$4.19 \pm 0.39$	$0.49 \pm 0.08$	$0.11 \pm 0.01$
21 <sub>4,18</sub> –20 <sub>3,17</sub>	727.379	252.1	$-5.61 \pm 0.11$	$2.39 \pm 0.26$	$0.71 \pm 0.15$	$0.30 \pm 0.03$
19 <sub>5,15</sub> –18 <sub>4,14</sub>	820.150	236.2	$-5.26 \pm 0.17$	$5.27 \pm 0.41$	$0.94 \pm 0.13$	$0.17 \pm 0.01$
20 <sub>5,15</sub> –19 <sub>4,16</sub>	840.751	254.6	$-4.79 \pm 0.20$	$5.25 \pm 0.49$	$0.70 \pm 0.11$	$0.12 \pm 0.01$
15 <sub>6,10</sub> –14 <sub>5,9</sub>	848.523	198.6	$-4.85 \pm 0.18$	$4.76 \pm 0.43$	$0.78 \pm 0.12$	$0.15 \pm 0.01$
<sup>34</sup> SO <sub>2</sub>						
19 <sub>1,19</sub> –18 <sub>0,18</sub> *	344.581	167.7	$-5.23 \pm 0.24$	$4.15 \pm 0.54$	$1.03 \pm 0.12$	$0.23 \pm 0.03$
13 <sub>4,10</sub> –13 <sub>3,11</sub> *	344.808	121.6	$-5.18 \pm 0.28$	$3.49 \pm 0.55$	$0.65 \pm 0.10$	$0.17 \pm 0.03$
7 <sub>4,4</sub> –7 <sub>3,5</sub> *	345.520	63.7	$-4.70 \pm 0.24$	$3.06 \pm 0.49$	$0.76 \pm 0.12$	$0.23 \pm 0.03$
20 <sub>0,20</sub> –19 <sub>1,19</sub> *	357.102	184.8	$-5.27 \pm 0.20$	$3.04 \pm 0.48$	$0.72 \pm 0.10$	$0.23 \pm 0.03$
6 <sub>3,3</sub> –5 <sub>2,4</sub> *	362.158	40.7	$-4.92 \pm 0.28$	$3.40 \pm 0.47$	$0.80 \pm 0.16$	$0.22 \pm 0.04$
o-H <sub>2</sub> O						
1 <sub>10</sub> –1 <sub>01</sub>	556.936	61.0	$-3.63 \pm 0.10$	$3.35 \pm 0.13$	$3.82 \pm 0.44$	$1.07 \pm 0.02$
			$-0.52 \pm 0.17$	$2.25 \pm 0.33$	$-0.66 \pm 0.42$	$-0.28 \pm 0.04$
3 <sub>12</sub> –3 <sub>03</sub>	1097.365	249.4	$-5.27 \pm 0.05$	$2.85 \pm 0.13$	$5.94 \pm 0.46$	$1.96 \pm 0.08$
			$-6.34 \pm 0.35$	$11.65 \pm 1.02$	$6.38 \pm 0.11$	$0.52 \pm 0.07$
3 <sub>12</sub> –2 <sub>21</sub>	1153.127	249.4	$-5.10 \pm 0.09$	$2.16 \pm 0.25$	$2.18 \pm 0.41$	$0.95 \pm 0.09$
			$-5.78 \pm 0.35$	$10.93 \pm 1.02$	$5.97 \pm 1.11$	$0.51 \pm 0.06$
3 <sub>21</sub> –3 <sub>12</sub>	1162.912	305.3	$-5.32 \pm 0.04$	$2.89 \pm 0.10$	$8.12 \pm 0.50$	$2.65 \pm 0.08$
			$-6.04 \pm 0.76$	$14.17 \pm 1.44$	$6.79 \pm 1.41$	$0.45 \pm 0.04$
p-H <sub>2</sub> O						
2 <sub>11</sub> –2 <sub>02</sub>	752.033	136.9	$-5.56 \pm 0.03$	$3.45 \pm 0.09$	$8.11 \pm 0.36$	$2.21 \pm 0.05$
			$-6.20 \pm 0.28$	$12.60 \pm 0.93$	$5.48 \pm 0.19$	$0.41 \pm 0.05$
2 <sub>02</sub> –1 <sub>11</sub>	987.927	100.9	$-5.33 \pm 0.04$	$3.82 \pm 0.10$	$11.12 \pm 0.45$	$2.74 \pm 0.06$
			$-5.81 \pm 0.40$	$14.78 \pm 1.30$	$8.01 \pm 0.19$	$0.54 \pm 0.05$
1 <sub>11</sub> –0 <sub>00</sub>	1113.343	53.4	$-3.30 \pm 0.20$	$3.26 \pm 0.34$	$6.17 \pm 0.63$	$1.78 \pm 0.11$
			$-0.48 \pm 0.20$	$2.60 \pm 0.27$	$-3.88 \pm 0.51$	$-1.40 \pm 0.17$
			$-11.98 \pm 0.34$	$13.75 \pm 0.84$	$-11.23 \pm 0.93$	$-0.77 \pm 0.03$
2 <sub>20</sub> –2 <sub>11</sub>	1228.789	195.9	$-5.24 \pm 0.10$	$2.82 \pm 0.27$	$6.23 \pm 0.64$	$2.08 \pm 0.20$
			$-5.98 \pm 0.91$	$8.47 \pm 1.59$	$5.66 \pm 1.80$	$0.63 \pm 0.20$
HF						
1–0	1232.476	59.2	$13.01 \pm 0.13$	$1.17 \pm 0.31$	$-1.16 \pm 0.52$	$-0.93 \pm 0.21$
			$-0.05 \pm 0.13$	$2.31 \pm 0.31$	$-2.81 \pm 0.76$	$-1.25 \pm 0.16$
			$-3.88 \pm 0.15$	$2.50 \pm 0.35$	$3.08 \pm 0.79$	$1.21 \pm 0.15$
			$-12.58 \pm 0.28$	$8.81 \pm 0.71$	$-10.02 \pm 1.39$	$-1.07 \pm 0.07$